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OM protein - protein search, using sw model

November 19, 2004, 16:34:27; Search time 19.3191 Seconds Run on:

(without alignments)

74.274 Million cell updates/sec

Title:

US-09-830-954A-1

Perfect score:

Sequence:

1 EFRH 4

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched:

2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters:

2002273

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

A Geneseq 23Sep04:\*

- 1: geneseqp1980s:\*
- 2: geneseqp1990s:\*
- 3: geneseqp2000s:\*
- 4: geneseqp2001s:\*
- 5: geneseqp2002s:\*
- 6: geneseqp2003as:\*
- 7: geneseqp2003bs:\*
- genesegp2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

R	esult		% Query						
	No.	Score		Length	DB	ID		Descript	ion
	1	24	100.0	4	2	AAW70870		Aaw70870	Beta-amyl
	2	24	100.0	4	6	AAO16062		Aao16062	Neurologi
	3	24	100.0	4	6	ABP70744		Abp70744	Antigenic
	4	24	100.0	4	7	ADB75167		Adb75167	Human amy
	5	24	100.0	4	7	ADE36574		Ade36574	Beta-amyl
	6	24	100.0	4	8	ADJ88108		Adj88108	Human bet
	7	24	100.0	4	8	ADJ71364		Adj71364	N-termina
	8	24	100.0	4	8	ADJ71377	•	Adj71377	N-termina
	9	24	100.0	4	8	ADP90808		Adp90808	Protein/p

24	100.0	5	6	ADA90172	Ada90172	Anti-Abet
24	100.0	5	8	ADJ71378	Adj71378	N-termina
24	100.0	5	8	ADJ71365	Adj71365	N-termina
24	100.0	5	8	ADJ71352	Adj71352	N-termina
24	100.0	6	2	AAW70868	Aaw70868	Beta-amyl
24	100.0	6	4	AAB47109	Aab47109	Epitope #
24	100.0	6	6	AA016067	Aao16067	Neurologi
24	100.0	6	6	ADA90170	Ada90170	Anti-Abet
24	100.0	6	7	ADB75165	Adb75165	Human amy
24	100.0	6	8	ADJ88114	Adj88114	fd phage
24	100.0	6	8	ADJ71366	Adj71366	N-termina
24	100.0	6	8	ADJ71379	Adj71379	N-termina
24	100.0	6	8	ADJ71340	Adj71340	N-termina
24	100.0	6	8	ADJ71353	Adj71353	N-termina
24	100.0	6	8	ADK52251	Adk52251	Human amy
24	100.0	6	8	ADK52264	Adk52264	Guinea pi
24	100.0	6	8	ADK52261	Adk52261	Rabbit am
24	100.0	6	8	ADK52260	Adk52260	Primate a
24	100.0	6	8	ADK52266	Adk52266	Amyloid b
24	100.0	7	4	AAB46202	Aab46202	Human APP
24	100.0	7	5	AA014421	Aao14421	Synthetic
24	100.0	7	6	AAO19884	Aao19884	Human amy
24	100.0	7	6	AAE35432	Aae35432	Abeta pep
24	100.0	7	6	ADA90925	Ada90925	Solid-pha
24	100.0	7	6 .	ADA90142	Ada90142	Anti-Abet
24	100.0	7	6	ADA90141	Ada90141	Anti-Abet
24	100.0	7	6	ADA90924	Ada90924	Solid-pha
24	100.0	7	6	ADA90171	Ada90171	Anti-Abet
24	100.0	7	8	ADJ71565	Adj71565	N-termina
24.	100.0	7	8	ADJ71380	Adj71380	N-termina
24	100.0	7	8	ADJ71341	Adj71341	N-termina
24	100.0	7	8	ADJ71367	Adj71367	N-termina
24	100.0	7		ADJ71354	Adj71354	N-termina
24	100.0	8	2	AAW70865	Aaw70865	Beta-amyl
24	100.0	8	5	AAU78518	Aau78518	N terminu
24	100.0	8	6	ABP70740	Abp70740	Antigenic
	24 24 24 24 24 24 24 24 24 24 24 24 24 2	24 100.0 24 100.0	24       100.0       5         24       100.0       5         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       6         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7         24       100.0       7      2	24       100.0       5       8         24       100.0       5       8         24       100.0       6       2         24       100.0       6       4         24       100.0       6       6         24       100.0       6       6         24       100.0       6       8         24       100.0       6       8         24       100.0       6       8         24       100.0       6       8         24       100.0       6       8         24       100.0       6       8         24       100.0       6       8         24       100.0       6       8         24       100.0       7       6         24       100.0       7       6         24       100.0       7       6         24       100.0       7       6         24       100.0       7       6         24       100.0       7       6         24       100.0       7       6         24       100.0       7       6	24 100.0 5 8 ADJ71378 24 100.0 5 8 ADJ71365 24 100.0 6 2 AAW70868 24 100.0 6 4 AAB47109 24 100.0 6 6 AAO16067 24 100.0 6 6 ADA90170 24 100.0 6 7 ADB75165 24 100.0 6 8 ADJ71336 24 100.0 6 8 ADJ71379 24 100.0 6 8 ADJ71353 24 100.0 6 8 ADJ88114 24 100.0 6 8 ADJ71353 24 100.0 6 8 ADK52251 24 100.0 6 8 ADK52264 24 100.0 6 8 ADK52261 24 100.0 6 8 ADK52266 24 100.0 6 8 ADK52266 24 100.0 7 4 AAB46202 24 100.0 7 6 AAO19884 24 100.0 7 6 AAO19884 24 100.0 7 6 ADA90142 24 100.0 7 6 ADA90142 24 100.0 7 6 ADA90141 24 100.0 7 6 ADA90141 24 100.0 7 6 ADA90171 24 100.0 7 8 ADJ71380 24 100.0 7 8 ADJ71380 24 100.0 7 8 ADJ71380 24 100.0 7 8 ADJ71367 24 100.0 7 8 ADJ71354	24         100.0         5         8         ADJ71378         Adj71378           24         100.0         5         8         ADJ71365         Adj71365           24         100.0         6         2         AAW70868         Aaw70868           24         100.0         6         4         AAB47109         Aab47109           24         100.0         6         6         AAD16067         Aac16067           24         100.0         6         6         AAD490170         Ada90170           24         100.0         6         7         ADB75165         Adb75165           24         100.0         6         8         ADJ71366         Adj71366           24         100.0         6         8         ADJ71379         Adj71379           24         100.0         6         8         ADJ71353         Adj71379           24         100.0         6         8         ADK52251         Adk52251           24         100.0         6         8         ADK52264         Adk52261           24         100.0         6         8         ADK52264         Adk52266           24         100.0         7 </td

## ALIGNMENTS

```
RESULT 1
AAW70870
     AAW70870 standard; peptide; 4 AA.
ΙD
XX
AC
     AAW70870;
XX
     04-FEB-1999
                  (first entry)
DT
XX
DE
     Beta-amyloid peptide epitope.
XX
     Beta-amyloid precursor protein; beta-APP; beta-amyloid peptide; antibody;
KW
     amyloid deposit; Alzheimer's disease.
KW
XX
     Synthetic.
OS
OS
     Homo sapiens.
XX
```

```
PN
     WO9844955-A1.
XX
PD
     15-OCT-1998.
XX
     09-APR-1998;
                    98WO-US006900.
ΡF
XX
PR
     09-APR-1997;
                    97US-0041850P.
XX
PA
     (MIND-) MINDSET LTD.
     (MCIN/) MCINNIS P A.
PA
XX
PI
     Chain DG;
XX
DR
     WPI; 1998-594476/50.
XX
     Preventing or inhibiting progression of Alzheimer's Disease - comprises
PT
РΤ
     use of recombinant DNA encoding an antibody specific for the N- or C-
РΤ
     terminus of an amyloid-beta peptide.
XX
     Example 1; Page 47; 58pp; English.
PS
XX
CC
     The present sequence represents a peptide epitope derived from beta-
     amyloid precursor protein peptide. The specification describes a method
CC
CC
     for prevention or inhibition of progression of Alzheimer's disease. The
     method comprises administering a composition comprising a recombinant DNA
CC
     molecule containing a gene encoding a recombinant antibody end-specific
CC
     for the N-terminus or the C-terminus of an amyloid-beta peptide, operably
CC
     linked to a promoter which is expressed in the central nervous system.
CC
     The recombinant antibody molecules prevent the accumulation of beta-
CC
     amyloid peptides in the extracellular space, interstitial fluid and
CC
     cerebrospinal fluid and the aggregation of such peptides into amyloid
CC
     deposits in the brain. They also inhibit the progression of Alzheimer's
CC
     disease by inhibiting the interaction of beta-amyloid peptides mediating
CC
     Alzheimer's disease induced neurotoxicity and inhibiting the Alzheimer's
CC
     disease induced complement activation and cytokine release involved in
CC
CC
     the inflammatory process
XX
SO
     Sequence 4 AA;
                          100.0%; Score 24; DB 2; Length 4;
  Query Match
  Best Local Similarity
                          100.0%; Pred. No. 1.7e+06;
                                                                  0; Gaps
  Matches
             4; Conservative
                                0; Mismatches
                                                   0: Indels
            1 EFRH 4
Qу
              \Pi\Pi\Pi
            1 EFRH 4
Dh
RESULT 2
AA016062
     AAO16062 standard; peptide; 4 AA.
TD
XX
AC
     AA016062;
XX
DT
     27-FEB-2003 (first entry)
XX
     Neurological/CNS disease treatment method-related peptide #1.
DE
```

```
XX
     Vaccine; gene therapy; neurological disease; CNS disorder;
KW
     central nervous system disorder; olfactory system; Alzheimer's disease;
ΚW
     Creutzfeld-Jakob disease; Huntington's chorea; Parkinson's disease;
KW
     viral infection of the brain; brain tumour; lysosomal storage disease;
KW
     multiple sclerosis.
KW
XX
OS
     Unidentified.
XX
     W0200274243-A2.
PN
XX
PD
     26-SEP-2002.
XX
PF
     15-MAR-2002; 2002WO-US008042.
XX
PR
     15-MAR-2001; 2001US-00808037.
XX
     (UYRA-) UNIV RAMOT APPLIED RES & IND DEV LTD.
PΑ
PΑ
     (MCIN/) MCINNIS P.
XX
     Solomon B, Frenkel D;
PΙ
XX
     WPI; 2003-040542/03.
DR
XX
     Treating or diagnosing neurological diseases of the central nervous
PT
     system, e.g. Alzheimer's disease, comprises displaying a polypeptide or
PT
     diagnostic agent on viral display vehicle and introducing or detecting
PT
PT
     the display vehicle.
XX
     Example 9; Page 138; 214pp; English.
PS
XX
     The invention comprises a method for treating a neurological disease or a
CC
     central nervous system (CNS) disorder. The method involves displaying a
CC
     therapeutic molecule capable of treating the neurological disease or CNS
CC
     disorder on a viral display vehicle. The viral display vehicle is then
CC
     introduced into the olfactory system of a subject to treat the disease or
CC
     disorder. The method of the invention is useful for preventing, treating
CC
     and diagnosing neurological diseases or CNS disorders, such as:
CC
     Alzheimer's disease; Creutzfeld-Jakob disease; Huntington's chorea; viral
CC
     infections of the brain; brain tumours; lysosomal storage diseases;
CC
     Parkinson's disease; and multiple sclerosis. The present amino acid
CC
     sequence represents a peptide which was used in the invention
CC
XX
SO
     Sequence 4 AA;
                           100.0%;
                                    Score 24; DB 6; Length 4;
  Query Match
                          100.0%;
  Best Local Similarity
                                    Pred. No. 1.7e+06;
                                  0; Mismatches
                                                                               0;
             4; Conservative
                                                    0; Indels
                                                                   0; Gaps
  Matches
            1 EFRH 4
Qу
               I \mid I \mid I
Db
            1 EFRH 4
RESULT 3
```

RESULT 3
ABP70744

ID ABP70744 standard; peptide; 4 AA.

```
XX
    ABP70744;
AC
XX
     15-MAY-2003 (first entry)
DT
XX
DE
     Antigenic peptide, SEQ ID 5.
XX
ΚW
     Nootropic; neuroprotective; antiinflammatory; vaccine; antigenic product;
KW
     plaque-forming disease; Alzheimer's disease; SAA amyloidosis;
     hereditary Icelandic syndrome; senility; multiple myeloma;
ΚW
     Creutzfeldt-Jakob disease; Kuru; Gerstmann-Straussler-Scheinker disease;
KW
     fatal familial insomnia; scrapie; bovine spongiform encephalitis;
KW
KW
     antigenic; multiantigen.
XX
OS
     Synthetic.
XX
PN
     WO2003000719-A2.
XX
PD
     03-JAN-2003.
XX
     20-JUN-2002; 2002WO-US019567.
PF
XX
PR
     20-JUN-2001; 2001US-0299201P.
PR
     12-APR-2002; 2002US-0371717P.
XX
PA
     (UYRA-) UNIV RAMOT.
PA
     (MCIN/) MCINNIS P.
XX
PΤ
     Mcinnis P, Solomon B;
XX
DR
     WPI; 2003-239139/23.
XX
     Antigenic product has dendritic polymer built on core molecule having
PΤ
     terminal functional groups to which antigenic peptide that has epitope of
PT
     deposit-forming polypeptide involved in plaque-forming disease is joined.
PT
XX
PS
     Claim 6; Page 44; 70pp; English.
XX
     The present invention relates to antigenic products (A), comprising a
CC
     dendritic polymer built on a core molecule which is at least difunctional
CC
     to provide branching and containing up to 16 terminal functional groups
CC
CC
     to which an antigenic peptide, that comprises an epitope of a deposit-
     forming polypeptide involved in plaque-forming disease, is joined by
CC
     covalent bonds. The antigenic products are useful for eliciting an immune
CC
     response against a deposit-forming polypeptide involved in a plaque-
CC
     forming disease or disorder, e.g. Alzheimer's disease, SAA amyloidosis,
CC
     hereditary Icelandic syndrome, senility, multiple myeloma, Creutzfeldt-
CC
     Jakob disease, Kuru, Gerstmann-Straussler-Scheinker disease, fatal
CC
CC
     familial insomnia, scrapie or bovine spongiform encephalitis, by
     administering the antigenic product to a subject in need of it. The
CC
     present sequence is one such antigenic peptide, which was used to
CC
CC.
     illustrate the invention
XX
SQ
     Sequence 4 AA;
                                   Score 24; DB 6; Length 4;
                          100.0%;
  Query Match
  Best Local Similarity 100.0%; Pred. No. 1.7e+06;
```

```
Matches
                 Conservative
                                                                  0; Gaps
                                  0; Mismatches
                                                    0; Indels
                                                                               0;
            1 EFRH 4
Qу
              1 EFRH 4
Db
RESULT 4
ADB75167
     ADB75167 standard; peptide; 4 AA.
TD
XX
AC
     ADB75167;
XX
DT
     04-DEC-2003 (first entry)
XX
DE
     Human amyloid beta peptide SEQ ID NO:8.
XX
KW
     antibody; amyloid beta peptide; amyloid beta; nootropic; neuroprotective;
KW
     antibody therapy; Alzheimer's disease; mild cognitive impairment;
     cerebral amyloid angiopathy; congiophylic angiopathy; Down's syndrome;
KW
     inclusion body myositis; neurotoxicity; beta amyloid precursor protein;
KW
     APP; human.
KW
XX
OS
     Homo sapiens.
XX
     WO2003074081-A1.
PΝ
XX
PD
     12-SEP-2003.
XX
PF
     21-OCT-2002; 2002WO-US031590.
XX
PR
     28-FEB-2002; 2002US-00084380.
XX
PA
     (MIND-) MINDSET BIOPHARMACEUTICALS USA INC.
XX
PΙ
     Chain DG;
XX
DR
     WPI; 2003-731651/69.
XX
PT
     New antibody that is targeted to amyloid beta peptide, or its fragment,
     useful for treating a subject having Alzheimer's disease, or a disease or
PT
PT
     disorder characterized by amyloid beta deposition, e.g. cognitive
PT
     impairment or dementia.
XX
     Disclosure; Page 60; 63pp; English.
PS
XX
CC
     The present invention describes an antibody that is targeted to amyloid
CC
     beta peptide, or its fragment. Also described: (1) an antibody that is
CC
     free-end specific and is targeted to: (a) the free N-terminus of amyloid
CC
     beta-peptide; (b) the free N-terminus of amyloid beta-peptide, where the
CC
     first amino acid of amyloid beta-peptide is aspartate; (c) the free N-
CC
     terminus of N- and/or C-terminus-truncated amyloid beta-peptide fragment;
CC
     (d) the free C-terminus of the amyloid beta-peptide Abetal-39, Abetal-40,
CC
     Abetal-41 or Abetal-43; or (e) to the free C-terminus of N- and/or C-
CC
     terminus-truncated amyloid beta-peptide fragment; (2) a single chain or
CC
     artificial antibody that is free-end specific and is targeted to the free
CC
     C-terminus of the amyloid beta-peptide Abeta1-42; and (3) a
```

```
CC
     pharmaceutical composition comprising the antibody, and a carrier. The
CC
     antibody targeted to amyloid beta peptide has nootropic and
CC
     neuroprotective activities, and can be used in antibody therapy. The
     antibody or its fragment is useful for manufacturing a medicament for
CC
     treating a subject having Alzheimer's disease, or a disease or disorder
CC
CC
     characterised by amyloid beta deposition (e.g. mild cognitive impairment,
CC
     cerebral amyloid angiopathy or congiophylic angiopathy, Alzheimer's
     disease associated with Down's syndrome, or inclusion body myositis), or
CC
CC
     for delaying, inhibiting or suppressing accumulation of amyloid beta
     peptide, or the neurotoxicity of amyloid beta peptide or its fragment.
CC
CC
     Amyloid beta peptide are derived from beta amyloid precursor protein
CC
     (APP). The present sequence represents an amyloid beta peptide which is
CC
     used in the exemplification of the present invention.
XX
SQ
     Sequence 4 AA;
 Query Match
                          100.0%; Score 24; DB 7; Length 4;
  Best Local Similarity 100.0%; Pred. No. 1.7e+06;
             4; Conservative
 Matches
                               0; Mismatches
                                                  0; Indels
                                                                  0; Gaps
                                                                              0;
            1 EFRH 4
Qy
              \mathbf{H}
Db
            1 EFRH 4
RESULT 5
ADE36574
     ADE36574 standard; peptide; 4 AA.
ID
XX
AC
     ADE36574;
XX
DT
     29-JAN-2004 (first entry)
XX
DE
     Beta-amyloid (Abeta) peptide 3-6 SEQ ID NO:2.
XX
KW
     immune response; beta-secretase cleavage site; amyloid precursor protein;
KW
     APP; nootropic; neuroprotective; vaccine; passive immunisation;
     Alzheimer's disease.
KW
XX
OS
     Synthetic.
XX
ΡN
     WO2003076455-A2.
XX
PD
     18-SEP-2003.
XX
PF
     04-MAR-2003; 2003WO-US006388.
XX
PR
     05-MAR-2002; 2002US-0361344P.
XX
PA
     (UYRA-) UNIV RAMOT AT TEL AVIV LTD.
PΑ
     (MCIN/) MCINNIS P.
XX
PΙ
     Solomon B;
XX
DR
     WPI; 2003-865017/80.
XX
PT
     Immunizing composition, useful for treating Alzheimer's disease by
```

```
inhibiting processing of amyloid precursor protein, also antibodies for
PT
     passive immunization.
XX
PS
     Disclosure; SEQ ID NO 2; 76pp; English.
XX
CC
     The present invention describes an immunising composition (A) comprising:
CC
     (a) an antigenic product (I) which induces an immune response against the
CC
     beta-secretase cleavage site of amyloid precursor protein (APP); and (b)
CC
     a carrier, diluent, excipient, adjuvant or auxiliary. Also described: (1)
CC
     a molecule (II) comprising the antigen-binding part of an antibody (Ab)
CC
     directed against the beta-secretase cleavage site of APP; (2) a
CC
     filamentous bacteriophage (FB) that displays (II), where this is a single
CC
     -chain Ab, on its surface; and (3) a composition containing FB. (A) has
CC
     nootropic and neuroprotective activities, and can be used in vaccines or
CC
     passive immunisation. (A) inhibits the cleavage of APP and so prevents
CC
     the formation of beta-amyloid. (A) can be used to induce an immune
CC
     response against the beta-secretase cleavage site of APP, specifically
CC
     for treatment and prevention of Alzheimer's disease. The molecule (II)
CC
     that contains the antigen-binding part of an Ab directed against the
CC
     cleavage site, or a filamentous phage that displays such an Ab (as a
CC
     single-chain molecule) can be used similarly, for passive immunisation.
CC
    The present sequence represents a beta-amyloid (Abeta) peptide which is
CC
     used in the exemplification of the present invention.
XX
SQ
     Sequence 4 AA;
  Query Match
                          100.0%;
                                   Score 24; DB 7; Length 4;
  Best Local Similarity
                          100.0%;
                                   Pred. No. 1.7e+06;
                               0; Mismatches
 Matches
            4; Conservative
                                                   0; Indels
                                                                  0; Gaps
                                                                              0;
Qy
            1 EFRH 4
              1111
Db
            1 EFRH 4
RESULT 6
ADJ88108
     ADJ88108 standard; peptide; 4 AA.
TD
XX
AC
    ADJ88108;
XX
DT
     06-MAY-2004 (first entry)
XX
DE
     Human beta amyloid peptide anti-agrregating epitope.
XX
KW
    Neurological disease; central nervous system; CNS disorder;
KW
     plaque-forming disease; Alzheimer's disease; SAA amyloidosis;
     hereditary Icelandic syndrome; senility; multiple myeloma; scrapie;
KW
KW
     bovine spongiform encephalopathy; BSE; kuru; Creutzfeldt-Jakob disease;
KW
     CJD; Gerstmann-Streussler-Sheinker disease; GSS; fatal familial insomnia;
     FFI; non-plaque-forming disease; Huntington's chorea; viral infection;
KW
     brain tumour; lysosomal storage disease; neurodegeneration;
KW
KW
     multiple sclerosis; vaccine; beta amyloid peptide; epitope; beta AP;
KW
     human.
XX
OS
     Homo sapiens.
XX
```

```
US2004013647-A1.
ΡN
XX
PD
     22-JAN-2004.
XX
ΡF
     11-MAR-2003; 2003US-00384788.
XX
PR
     03-SEP-1999;
                    99US-0152417P.
PR
     29-DEC-1999:
                    99US-00473653.
     31-JUL-2000; 2000US-00629971.
PR
     31-AUG-2000; 2000WO-IL000518.
PR
PR
     15-MAR-2001; 2001US-00808037.
PR
     07-AUG-2001; 2001US-00830954.
PR
     12-APR-2002; 2002US-0371735P.
PR
     06-JUN-2002; 2002US-00162889.
XX
PΑ
     (UYRA-) UNIV RAMOT AT TEL AVIV LTD.
XX
PΙ
     Solomon B, Frenkel D;
XX
DR
     WPI; 2004-108188/11.
XX
     Treating neurological disease CNS e.g., Alzheimer's disease, by
PT
     displaying therapeutic molecule capable of treating the disease on viral
PT
     display vehicle which is then administered to subject through olfactory
PT
PT
     system.
XX
     Example 10; SEQ ID NO 1; 68pp; English.
PS
XX
     The invention relates to a method of treating a neurological disease or
CC
     disorder of the central nervous system (CNS). The method involves
CC
     displaying a therapeutic molecule capable of treating the neurological
CC
     disease or disorder of the CNS on a viral display vehicle and introducing
CC
     viral display vehicle into a subject by applying an effective amount of
CC
     the viral display vehicle displaying the therapeutic molecule to an
CC
     olfactory system of the subject. The method is useful for treating a
CC
     neurological disease or disorder of CNS such as a plaque-forming disease
CC
     such as Alzheimer's disease, late onset Alzheimer's disease,
CC
     presymptomatic Alzheimer's disease, SAA amyloidosis, hereditary Icelandic
CC
     syndrome, senility, multiple myeloma, scrapie, bovine spongiform
CC
     encephalopathy (BSE), kuru, Creutzfeldt-Jakob disease (CJD), Gerstmann-
CC
     Streussler-Sheinker disease (GSS) or fatal familial insomnia (FFI). The
CC
CC
     method is also useful for treating a non plaque forming disease or
CC
     disorder e.g. Huntington's chorea, viral infections of the brain, brain
CC
     tumours, lysosomal storage diseases which cause neurodegeneration and are
     manifested by enzyme deficiencies and multiple sclerosis. The invention
CC
     is also used in the preparation of vaccines. The present sequence is
CC
     human beta amyloid peptide (beta AP) anti-agrregating epitope. This
CC
     sequence is used to illustrate the method of the invention.
CC
XX
SQ
     Sequence 4 AA;
                          100.0%;
                                   Score 24; DB 8; Length 4;
  Query Match
                          100.0%;
                                   Pred. No. 1.7e+06;
  Best Local Similarity
                                 0; Mismatches
                                                                              0;
                                                                  0; Gaps
             4; Conservative
                                                    0; Indels
  Matches
            1 EFRH 4
```

++++

1111

```
RESULT 7
ADJ71364
     ADJ71364 standard; peptide; 4 AA.
ID
XX
AC
     ADJ71364;
XX
     06-MAY-2004 (first entry)
DT
XX
DΕ
     N-terminal truncated beta-amyloid peptide, SEQ ID 27.
XX
ΚW
     Nootropic; Neuroprotective; Vaccine; beta Amyloid;
KW
     amyloid precursor protein; APP; Alzheimer's disease.
XX
os
     Homo sapiens.
XX
PN
     WO2004013172-A2.
XX
PD
     12-FEB-2004.
XX
PF
     18-JUL-2003; 2003WO-EP007833.
XX
PR
     24-JUL-2002; 2002EP-00447147.
     06-AUG-2002; 2002US-0401497P.
PR
XX
PA
     (INNO-) INNOGENETICS NV.
XX
PΙ
     Delacourte A, Sergeant N;
XX
DR
     WPI; 2004-180423/17.
XX
PT
     New beta-amyloid or amyloid precursor protein preparation, useful as a
     prophylactic vaccine or a therapeutic for preventing or treating a
PT
     disease associated with beta-amyloid formation and/or aggregation, e.g.
PT
PT
     Alzheimer's disease.
XX
PS
     Claim 4; Page 61; 104pp; English.
XX
CC
     The present invention relates to preparations (I) comprising a beta-
CC
     amyloid peptide variant or beta-amyloid N-terminal fragment, or N-
     terminal amyloid precursor protein (APP) soluble fragment or C-terminal
CC
CC
     fragment. The beta-amyloid or APP preparations are useful for
CC
     manufacturing a prophylactic vaccine or a therapeutic, or as a
CC
     prophylactic vaccine for the prevention, or as a therapeutic for the
CC
     treatment of a disease associated with beta-amyloid formation and/or
CC
     aggregation, such as Alzheimer's disease.
XX
SQ
     Sequence 4 AA;
  Query Match
                           100.0%;
                                   Score 24; DB 8; Length 4;
                                    Pred. No. 1.7e+06;
  Best Local Similarity
                          100.0%;
                                  0; Mismatches
  Matches
             4; Conservative
                                                                  0; Gaps
                                                                               0;
                                                    0; Indels
            1 EFRH 4
Ov
```

```
RESULT 8
ADJ71377
     ADJ71377 standard; peptide; 4 AA.
ID
XX
AC
     ADJ71377;
XX
DT
     06-MAY-2004 (first entry)
XX
DΕ
     N-terminal truncated beta-amyloid peptide, SEQ ID 40.
XX
KW
     Nootropic; Neuroprotective; Vaccine; beta Amyloid;
KW
     amyloid precursor protein; APP; Alzheimer's disease.
XX
     Homo sapiens.
OS
XX
FΗ
     Key
                     Location/Qualifiers
FT
     Modified-site
                     /note= "Pyroglutamic acid"
FT
XX
PN
     WO2004013172-A2.
XX
PD
     12-FEB-2004.
XX
PF
     18-JUL-2003; 2003WO-EP007833.
XX
PR
     24-JUL-2002; 2002EP-00447147.
     06-AUG-2002; 2002US-0401497P.
PR
XX
     (INNO-) INNOGENETICS NV.
PA
XX
PI
     Delacourte A, Sergeant N;
XX
     WPI: 2004-180423/17.
DR
XX
     New beta-amyloid or amyloid precursor protein preparation, useful as a
PT
PT
     prophylactic vaccine or a therapeutic for preventing or treating a
PT
     disease associated with beta-amyloid formation and/or aggregation, e.g.
PT
     Alzheimer's disease.
XX
     Claim 4; Page 62; 104pp; English.
PS
XX
     The present invention relates to preparations (I) comprising a beta-
CC
     amyloid peptide variant or beta-amyloid N-terminal fragment, or N-
CC
     terminal amyloid precursor protein (APP) soluble fragment or C-terminal
CC
     fragment. The beta-amyloid or APP preparations are useful for
CC
CC
     manufacturing a prophylactic vaccine or a therapeutic, or as a
     prophylactic vaccine for the prevention, or as a therapeutic for the
CC
     treatment of a disease associated with beta-amyloid formation and/or
CC
     aggregation, such as Alzheimer's disease.
CC
XX
     Sequence 4 AA;
SO
                           100.0%; Score 24; DB 8; Length 4;
  Query Match
  Best Local Similarity 100.0%; Pred. No. 1.7e+06;
```

```
Conservative
                                 0; Mismatches
                                                   0; Indels
                                                                  0; Gaps
                                                                              0;
  Matches
            1 EFRH 4
Qу
              1 EFRH 4
Db
RESULT 9
ADP90808
     ADP90808 standard; peptide; 4 AA.
ID
XX
AC
     ADP90808;
XX
     09-SEP-2004 (first entry)
DT
XX
     Protein/peptide labelling method-related affinity tag peptide #3.
DΕ
XX
KW
     protein labelling; peptide labelling;
KW
     irreversible affinity tagging residue;
     reversible affinity tagging residue; high throughput screening assay;
KW
     pharmaceutical agent; affinity tag.
KW
XX
     Unidentified.
OS
XX
                     Location/Qualifiers
FH
     Key
FT
     Modified-site
                     /note= "C-terminal amide"
FT
XX
     WO2004051270-A2.
PN
XX
     17-JUN-2004.
PD
XX
     04-DEC-2003; 2003WO-EP013715.
PF
XX
PR
     05-DEC-2002; 2002GB-00028429.
XX
PA
     (NOVS ) NOVARTIS AG.
     (NOVS ) NOVARTIS PHARMA GMBH.
PA
XX
     Auer M, Meisner N, Seifert J;
PI
XX
     WPI; 2004-480677/45.
DR
XX
     Providing labeled target protein or target peptide by contacting chemical
PT
     compound with affinity support, removing impurities in reaction mixture
PT
PT
     surrounding affinity support, cleaving or eluting chemical molecule from
PT
     affinity support.
XX
     Claim 4; Page 70; 81pp; English.
PS
XX
CC
     The invention comprises a method for providing a labelled target
CC
     protein/peptide. The method involves contacting a chemical compound with
     affinity support, removing impurities in the reaction mixture surrounding
CC
CC
     the affinity support to which the chemical molecule is bound, and
     cleaving or eluting the molecule from the affinity support to obtain
CC
     irreversible or reversible affinity tagging residue, labelled target
CC
     protein or labelled peptide. The method of the invention is useful for
CC
```

```
CC
     labelling a target protein/peptide or high throughput screening assay.
     The method of the invention is useful for identifying agents that
CC
     modulate the activity or characteristics of a target protein/peptide -
CC
CC
     such agents are useful as pharmaceuticals. The present amino acid
CC
     sequence represents an affinity tag peptide of the invention.
XX
     Sequence 4 AA;
SO
  Query Match
                          100.0%; Score 24; DB 8; Length 4;
                          100.0%; Pred. No. 1.7e+06;
  Best Local Similarity
             4; Conservative
                                0; Mismatches
                                                   0; Indels
                                                                  0; Gaps
                                                                              0;
            1 EFRH 4
·Qγ
              1111
Db
            1 EFRH 4
RESULT 10
ADA90172
     ADA90172 standard; peptide; 5 AA.
XX
АC
     ADA90172;
XX
     20-NOV-2003 (first entry)
DT
XX
DE
     Anti-Abeta antibody related amino acid sequence SEQ ID NO:287.
XX
KW
     antibody molecule; antibody; beta-A4 peptide; Abeta4; neuroprotective;
KW
     nootropic; antiparkinsonian; gene therapy; amyloidogenesis;
KW
     amyloid-plaque formation; beta-amyloid plaque; immunisation; dementia;
KW
     Alzheimer's disease; motor neuropathy; Down's syndrome;
     Creutzfeldt Jacob disease; hereditary cerebral haemorrhage; amyloidosis;
KW
     Parkinson's disease; HIV-related dementia; amyotrophic lateral sclerosis;
KW
KW
     neuronal disorder; aging.
XX
OS
     Synthetic.
OS
     Homo sapiens.
XX
PN
     WO2003070760-A2.
XX
PD
     28-AUG-2003.
XX
     20-FEB-2003; 2003WO-EP001759.
PF
XX
     20-FEB-2002; 2002EP-00003844.
PR
XX
PA
     (HOFF ) HOFFMANN LA ROCHE & CO AG F.
PA
     (MORP-) MORPHOSYS AG.
XX
                  Bohrmann B, Brockhaus M, Huber W, Kretzschmar T;
PI
     Bardroff M,
PI
     Loehning C, Loetscher H, Nordstedt C, Rothe C;
XX
DR
     WPI; 2003-663848/62.
XX
     New antibody molecule capable of specifically recognizing two regions of
PT
     the beta-A4 peptide, useful for diagnosing, preventing or treating
PT
     diseases associated with amyloidogenesis or amyloid-plaque formation
PT
```

```
PT
     (e.g. dementia).
XX
PS
     Disclosure; Page 265; 312pp; English.
XX
CC
     The present invention describes an antibody molecule (I) capable of
CC
     specifically recognising two regions of the beta-A4 peptide/Abeta4. The
CC
     first region comprises the amino acid sequence Ala-Glu-Phe-Arg-His-Asp-
CC
     Ser-Gly-Tyr ADA89886 or its fragment, and the second region comprises the
     amino acid sequence Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-
CC
CC
     Gly ADA89887 or its fragment. Also described: (1) a nucleic acid molecule
CC
     encoding (I); (2) a vector comprising the nucleic acid of (1); (3) a host
CC
     cell comprising the vector of (2); (4) preparing (I), comprising
CC
     culturing the host cell of (3) under conditions that allow synthesis of
CC
     (I) and recovering (I) from the culture; (5) a composition comprising (I)
CC
     or an antibody molecule produced by method (4); (6) a kit comprising (I),
     nucleic acid of (1), vector of (2) or host cell of (3); (7) optimising
CC
     (I); (8) testing the resulting Fab optimisation library by panning
CC
CC
     against Abeta/Abeta4; (9) identifying optimised clones; (10) expressing
CC
     of selected, optimised clones; (11) preparing a pharmaceutical
CC
     composition, comprising optimisation of (I), and formulating the
CC
     optimised antibody/antibody molecule with a carrier; and (12) a
CC
     pharmaceutical composition prepared by method (8). (I) has
CC
     neuroprotective, nootropic and antiparkinsonian activities, and can be
CC
     used in gene therapy. The antibody molecule (I), nucleic acid molecule,
CC
     vector or host is useful in preparing a pharmaceutical composition for
CC
     the prevention and/or treatment of a disease associated with
CC
     amyloidogenesis and/or amyloid-plaque formation. The antibody molecule
CC
     may also be used in preparing a diagnostic composition for the detection
CC
     of the disease mentioned above. The antibody is used for the
CC
     disintegration of beta-amyloid plaques or for passive immunisation
CC
     against beta-amyloid plaque formation. In particular, the disease is
CC
     dementia, Alzheimer's disease, motor neuropathy, Down's syndrome,
CC
     Creutzfeldt Jacob disease, hereditary cerebral haemorrhage with
CC
     amyloidosis Dutch type, Parkinson's disease, HIV-related dementia,
CC
     amyotrophic lateral sclerosis or neuronal disorders related to aging. The
CC
     present sequence is used in the exemplification of the present invention.
XX
SQ
     Sequence 5 AA;
  Query Match
                          100.0%; Score 24; DB 6; Length 5;
  Best Local Similarity
                          100.0%; Pred. No. 1.7e+06;
  Matches
            4; Conservative
                                0; Mismatches
                                                   0;
                                                       Indels
                                                                      Gaps
                                                                              0;
            1 EFRH 4
Qу
              \mathbf{H}
Db
            1 EFRH 4
RESULT 11
ADJ71378
     ADJ71378 standard; peptide; 5 AA.
TD
XX
AC
     ADJ71378;
XX
DT
     06-MAY-2004 (first entry)
XX
DE
     N-terminal truncated beta-amyloid peptide, SEQ ID 41.
```

```
XX
     Nootropic; Neuroprotective; Vaccine; beta Amyloid;
KW
     amyloid precursor protein; APP; Alzheimer's disease.
KW
XX
os
     Homo sapiens.
XX
                     Location/Qualifiers
FH
     Key
     Modified-site
FT
                     /note= "Pyroglutamic acid"
FΤ
XX
     WO2004013172-A2.
PN
XX
     12-FEB-2004.
PD
XX
     18-JUL-2003; 2003WO-EP007833.
PF
XX
     24-JUL-2002; 2002EP-00447147.
PR
     06-AUG-2002; 2002US-0401497P.
PR
XX
     (INNO-) INNOGENETICS NV.
PΑ
XX
     Delacourte A, Sergeant N;
PΙ
XX
     WPI; 2004-180423/17.
DR
XX
     New beta-amyloid or amyloid precursor protein preparation, useful as a
PT
     prophylactic vaccine or a therapeutic for preventing or treating a
PT
     disease associated with beta-amyloid formation and/or aggregation, e.g.
PT
     Alzheimer's disease.
PT
XX
     Claim 4; Page 62; 104pp; English.
PS
XX
     The present invention relates to preparations (I) comprising a beta-
CC
     amyloid peptide variant or beta-amyloid N-terminal fragment, or N-
CC
     terminal amyloid precursor protein (APP) soluble fragment or C-terminal
CC
     fragment. The beta-amyloid or APP preparations are useful for
CC
     manufacturing a prophylactic vaccine or a therapeutic, or as a
CC
     prophylactic vaccine for the prevention, or as a therapeutic for the
CC
     treatment of a disease associated with beta-amyloid formation and/or
CC
     aggregation, such as Alzheimer's disease.
CC
XX
 SO
      Sequence 5 AA;
                           100.0%; Score 24; DB 8; Length 5;
  Query Match
                           100.0%; Pred. No. 1.7e+06;
  Best Local Similarity
                                                                               0;
                                                                   0; Gaps
                                                    0; Indels
                                0; Mismatches
              4; Conservative
             1 EFRH 4
 Qу
               1111
             1 EFRH 4
 Db
 RESULT 12
 ADJ71365
      ADJ71365 standard; peptide; 5 AA.
 ID
 XX
 AC
      ADJ71365;
```

```
XX
DT
     06-MAY-2004 (first entry)
XX
     N-terminal truncated beta-amyloid peptide, SEQ ID 28.
DΕ
XX
     Nootropic; Neuroprotective; Vaccine; beta Amyloid;
KW
     amyloid precursor protein; APP; Alzheimer's disease.
KW
XX
OS
     Homo sapiens.
XX
PN
     WO2004013172-A2.
XX
PD
     12-FEB-2004.
XX
PF
     18-JUL-2003; 2003WO-EP007833.
XX
     24-JUL-2002; 2002EP-00447147.
PR
     06-AUG-2002; 2002US-0401497P.
PR
XX
     (INNO-) INNOGENETICS NV.
PΑ
XX
PI
     Delacourte A, Sergeant N;
XX
     WPI; 2004-180423/17.
DR
XX
     New beta-amyloid or amyloid precursor protein preparation, useful as a
PT
     prophylactic vaccine or a therapeutic for preventing or treating a
PT
     disease associated with beta-amyloid formation and/or aggregation, e.g.
PT
     Alzheimer's disease.
PT
XX
     Claim 4; Page 61; 104pp; English.
PS
XX
     The present invention relates to preparations (I) comprising a beta-
CC
     amyloid peptide variant or beta-amyloid N-terminal fragment, or N-
CC
     terminal amyloid precursor protein (APP) soluble fragment or C-terminal
CC
     fragment. The beta-amyloid or APP preparations are useful for
CC
СĆ
     manufacturing a prophylactic vaccine or a therapeutic, or as a
     prophylactic vaccine for the prevention, or as a therapeutic for the
CC
     treatment of a disease associated with beta-amyloid formation and/or
CC
CC
     aggregation, such as Alzheimer's disease.
XX
SQ
     Sequence 5 AA;
  Query Match
                          100.0%; Score 24; DB 8; Length 5;
                          100.0%;
                                   Pred. No. 1.7e+06;
  Best Local Similarity
             4; Conservative
                                                   0; Indels 0; Gaps
                               0; Mismatches
  Matches
            1 EFRH 4
Qу
              \mathbf{H}
            1 EFRH 4
Db
RESULT 13
ADJ71352
     ADJ71352 standard; peptide; 5 AA.
ΙD
XX
AC
     ADJ71352:
```

```
XX
     06-MAY-2004 (first entry)
DT
XX
     N-terminal truncated beta-amyloid peptide, SEQ ID 15.
DΕ
XX
     Nootropic; Neuroprotective; Vaccine; beta Amyloid;
KW
     amyloid precursor protein; APP; Alzheimer's disease.
KW
XX
OS
     Homo sapiens.
XX
FH
     Key
                     Location/Qualifiers
FT
     Modified-site
                     /note= "Optionally methylated"
FT
XX
PN
     WO2004013172-A2.
XX
     12-FEB-2004.
PD
ΧX
     18-JUL-2003; 2003WO-EP007833.
PF
XX
     24-JUL-2002; 2002EP-00447147.
PR
     06-AUG-2002; 2002US-0401497P.
PR
XX
     (INNO-) INNOGENETICS NV.
PΑ
XX
     Delacourte A, Sergeant N;
PI
XX
DR
     WPI; 2004-180423/17.
XX
     New beta-amyloid or amyloid precursor protein preparation, useful as a
PT
     prophylactic vaccine or a therapeutic for preventing or treating a
PT
     disease associated with beta-amyloid formation and/or aggregation, e.g.
     Alzheimer's disease.
PT
XX
     Claim 4; Page 61; 104pp; English.
PS
XX
     The present invention relates to preparations (I) comprising a beta-
CC
     amyloid peptide variant or beta-amyloid N-terminal fragment, or N-
CC
     terminal amyloid precursor protein (APP) soluble fragment or C-terminal
CC
     fragment. The beta-amyloid or APP preparations are useful for
CC
     manufacturing a prophylactic vaccine or a therapeutic, or as a
CC
     prophylactic vaccine for the prevention, or as a therapeutic for the
CC
     treatment of a disease associated with beta-amyloid formation and/or
CC
CC
     aggregation, such as Alzheimer's disease.
XX
     Sequence 5 AA;
SQ
                           100.0%; Score 24; DB 8; Length 5;
  Query Match
                           100.0%;
                                   Pred. No. 1.7e+06;
  Best Local Similarity
             4; Conservative
                                  0; Mismatches
                                                    0; Indels
                                                                  0;
                                                                      Gaps
                                                                               0;
  Matches
            1 EFRH 4
Qу
               1111
            2 EFRH 5
Db
```

```
AAW70868 standard; peptide; 6 AA.
ΙD
XX
AC
     AAW70868;
XX
DT
     04-FEB-1999 (first entry)
XX
DE
     Beta-amyloid peptide to create a monoclonal antibody.
XX
KW
     Beta-amyloid precursor protein; beta-APP; beta-amyloid peptide; antibody;
KW
     amyloid deposit; Alzheimer's disease.
XX
OS
     Synthetic.
OS
     Homo sapiens.
XX
PN
     WO9844955-A1.
XX
PD
     15-OCT-1998.
XX
     09-APR-1998;
                    98WO-US006900.
PF
XX
     09-APR-1997;
                    97US-0041850P.
PR
XX
     (MIND-) MINDSET LTD.
PA
     (MCIN/) MCINNIS P A.
PA
XX
PΙ
     Chain DG;
XX
DR
     WPI; 1998-594476/50.
XX
PT
     Preventing or inhibiting progression of Alzheimer's Disease - comprises
PT
     use of recombinant DNA encoding an antibody specific for the N- or C-
PΤ
     terminus of an amyloid-beta peptide.
XX
PS
     Example 1; Page 47; 58pp; English.
XX
CC
    The present sequence represents a peptide derived from beta-amyloid
     precursor protein (beta-APP, see AAW70863). The peptide is a beta-amyloid
CC
CC
     peptide and is used to produce a monoclonal antibody. The specification
CC
     describes a method for prevention or inhibition of progression of
CC
     Alzheimer's disease. The method comprises administering a composition
CC
     comprising a recombinant DNA molecule containing a gene encoding a
CC
     recombinant antibody end-specific for the N-terminus or the C-terminus of
CC
     an amyloid-beta peptide, operably linked to a promoter which is expressed
CC
     in the central nervous system. The recombinant antibody molecules prevent
     the accumulation of beta-amyloid peptides in the extracellular space,
CC
     interstitial fluid and cerebrospinal fluid and the aggregation of such
CC
CC
     peptides into amyloid deposits in the brain. They also inhibit the
     progression of Alzheimer's disease by inhibiting the interaction of beta-
CC
CC
     amyloid peptides mediating Alzheimer's disease induced neurotoxicity and
CC
     inhibiting the Alzheimer's disease induced complement activation and
     cytokine release involved in the inflammatory process
CC
XX
SQ
     Sequence 6 AA;
  Query Match
                          100.0%;
                                   Score 24; DB 2; Length 6;
  Best Local Similarity
                          100.0%; Pred. No. 1.7e+06;
```

```
0; Indels
 Matches
             4; Conservative
                                 0; Mismatches
                                                                  0;
                                                                      Gaps
                                                                              0;
            1 EFRH 4
Qу
              3 EFRH 6
Db
RESULT 15
AAB47109
     AAB47109 standard; peptide; 6 AA.
ID
XX
AC
     AAB47109;
XX
DT
     04-JUN-2001 (first entry)
XX
DΕ
     Epitope #1 used in treatment of plaque forming disease.
XX
     Human; prion protein; plaque forming disease; display vehicle; kuru;
KW
     aggregating protein; amyloid plaque; brain; early onset; senility;
KW
KW
     Alzheimer's disease; late onset; pre-symptomatic; SAA amyloidosis;
     hereditary Icelandic syndrome; multiple myeloma; scrapie; BSE; CJD;
KW
     bovine spongiform encephalopathy; Creutzfeldt-Jakob Disease; FFI;
KW
     Gerstmann-Streussler-Sheinker Disease; GSS; fatal familial insomnia.
KW
XX
OS
     Synthetic.
XX
     WO200118169-A2.
PN
XX
PD
     15-MAR-2001.
XX
PF
     31-AUG-2000; 2000WO-IL000518.
XX
PR
     03-SEP-1999;
                    99US-0152417P.
PR
     29-DEC-1999;
                    99US-00473653.
     31-JUL-2000; 2000US-00629971.
PR
XX
PΑ
     (UYRA-) UNIV RAMOT APPLIED RES & IND DEV LTD.
XX
PI
     Solomon B, Frenkel D, Hanan E;
XX
DR
     WPI; 2001-244564/25.
XX
PT
     Treating amyloidgenic disease such as Alzheimer's disease, BSE or CJD
     comprises presentation of plaque derived antiqens or epitopes on a
PT
PT
     display vehicle, and introducing the vehicle into the recipient.
XX
PS
     Example; Page 50; 120pp; English.
XX
CC
     This peptide is based on the N-terminal fragment of beta-amyloid peptide
     (beta-AP) and was fused to the minor coat protein of fd phage. This
CC
     peptide may be used in the method of the invention. The invention
CC
     provides an agent for treating a plaque forming disease. The polypeptide
CC
     is displayed on a display vehicle and is capable of eliciting antibodies
CC
     capable of disaggregating the aggregating protein and/or of preventing
СÇ
     aggregation of the aggregating protein. This reduces formation of amyloid
CC
     plaques in the brain of victims of plaque forming diseases, e.g. early
CC
CC
     onset Alzheimer's disease, late onset Alzheimer's disease, pre-
```

```
symptomatic Alzheimer's disease, SAA amyloidosis, hereditary Icelandic
CC
     syndrome, senility, multiple myeloma, scrapie, bovine spongiform
CC
     encephalopathy (BSE), kuru, Creutzfeldt-Jakob Disease (CJD), Gerstmann-
CC
     Streussler-Sheinker Disease (GSS) and fatal familial insomnia (FFI)
XX
SQ
     Sequence 6 AA;
  Query Match
                         100.0%; Score 24; DB 4; Length 6;
                         100.0%; Pred. No. 1.7e+06;
  Best Local Similarity
  Matches
           4; Conservative 0; Mismatches 0; Indels
                                                               0; Gaps
                                                                           0;
Qy.
           1 EFRH 4
             1111
           3 EFRH 6
Db .
```

Search completed: November 19, 2004, 16:54:07 Job time: 21.3191 secs

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OM protein - protein search, using sw model

November 19, 2004, 16:39:17; Search time 4.68085 Seconds Run on:

(without alignments)

56.672 Million cell updates/sec

Title:

US-09-830-954A-1

Perfect score:

2.4

Sequence:

1 EFRH 4

Scoring table:

BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched:

478139 seqs, 66318000 residues

Total number of hits satisfying chosen parameters:

478139

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

9

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents AA:\*

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- 2: /cgn2 6/ptodata/1/iaa/5B COMB.pep:\*
- 3: /cgn2 6/ptodata/1/iaa/6A\_COMB.pep:\*
- 4: /cgn2 6/ptodata/1/iaa/6B\_COMB.pep:\*
- 5: /cgn2 6/ptodata/1/iaa/PCTUS COMB.pep:\*
- 6: /cgn2 6/ptodata/1/iaa/backfiles1.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

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2	24	100.0	7	4	US-09-579-012-25	Sequence 25, Appl
3	24	100.0	10	1	US-08-371-930-12	Sequence 12, Appl
4	24	100.0	10	4	US-09-724-961-5	Sequence 5, Appli
5	24	100.0	10	4	US-09-724-961-6	Sequence 6, Appli
6	24	100.0	. 10	4	US-09-724-961-7	Sequence 7, Appli
7	24	100.0	10	4	US-09-724-961-8	Sequence 8, Appli
8	24	100.0	10	4	US-09-724-961-9	Sequence 9, Appli
9	24	100.0	10	4	US-09-724-961-10	Sequence 10, Appl
10	24	100.0	10	4	US-09-724-961-11	Sequence 11, Appl
11	24	100.0	10	4	US-09-580-018-5	Sequence 5, Appli

12	24	100.0	10	4	US-09-580-018-6	Sequence	6, Appli
13	24	100.0	10	4	US-09-580-018-7	Sequence	7, Appli
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## ALIGNMENTS

# RESULT 1

US-09-579-012-24

- ; Sequence 24, Application US/09579012
- ; Patent No. 6670195
- ; GENERAL INFORMATION:
- ; APPLICANT: Jorge GHISO
- ; APPLICANT: Ruben VIDAL
- ; APPLICANT: Blas FRANGIONE
- ; TITLE OF INVENTION: New Mutant Genes in Familial British Dementia and

## Familial Danish

- ; TITLE OF INVENTION: Dementia
- ; FILE REFERENCE: 32004-16277
- ; CURRENT APPLICATION NUMBER: US/09/579,012
- ; CURRENT FILING DATE: 2000-05-26
- ; PRIOR APPLICATION NUMBER: US 60/136238
- ; PRIOR FILING DATE: 1999-05-26
- ; NUMBER OF SEQ ID NOS: 28
- ; SOFTWARE: PatentIn version 3.1

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Qу
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; Sequence 25, Application US/09579012
; Patent No. 6670195
; GENERAL INFORMATION:
  APPLICANT: Jorge GHISO
 APPLICANT: Ruben VIDAL
; APPLICANT: Blas FRANGIONE
  TITLE OF INVENTION: New Mutant Genes in Familial British Dementia and
Familial Danish
; TITLE OF INVENTION: Dementia
; FILE REFERENCE: 32004-16277
  CURRENT APPLICATION NUMBER: US/09/579,012
  CURRENT FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/136238
; PRIOR FILING DATE: 1999-05-26
; NUMBER OF SEQ ID NOS: 28
; SOFTWARE: PatentIn version 3.1
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; Sequence 12, Application US/08371930
; Patent No. 5578451
  GENERAL INFORMATION:
     APPLICANT: Nishimoto, Ikuo
; TITLE OF INVENTION: ALZHEIMER'S DISEASE THERAPEUTICS
     NUMBER OF SEQUENCES: 30
     CORRESPONDENCE ADDRESS:
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ADDRESSEE: Fish & Richardson
      STREET: 225 Franklin Street
      CITY: Boston
      STATE: Massachusetts
      COUNTRY: U.S.A.
      ZIP: 02110-2804
    COMPUTER READABLE FORM:
      MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
      COMPUTER: IBM PS/2 Model 50Z or 55SX
      OPERATING SYSTEM: MS-DOS (Version 5.0)
      SOFTWARE: WordPerfect (Version 5.1)
    CURRENT APPLICATION DATA:
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      FILING DATE:
      CLASSIFICATION: 436
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 08/019,208
      FILING DATE: February 18, 1993
    ATTORNEY/AGENT INFORMATION:
      NAME: Clark, Paul T.
      REGISTRATION NUMBER: 30,162
      REFERENCE/DOCKET NUMBER: 00786/154001
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (617) 542-5070
      TELEFAX: (617) 542-8906
      TELEX: 200154
  INFORMATION FOR SEQ ID NO: 12:
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      TOPOLOGY: linear
US-08-371-930-12
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; Sequence 5, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
; APPLICANT: Schenk, Dale B.
 APPLICANT: Bard, Frederique
 APPLICANT: Vasquez, Nicki
 APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
  FILE REFERENCE: 15270J-004750UC
; CURRENT APPLICATION NUMBER: US/09/724,961
; CURRENT FILING DATE: 2000-11-28
; PRIOR APPLICATION NUMBER: US 09/580,015
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  PRIOR APPLICATION NUMBER: US 09/322,289
   PRIOR FILING DATE: 1999-05-28
   PRIOR APPLICATION NUMBER: US 09/201,430
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: WO PCT/US00/14810
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: US 60/080,970
   PRIOR FILING DATE: 1998-04-07
   PRIOR APPLICATION NUMBER: US 60/067,740
   PRIOR FILING DATE: 1997-12-02
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    OTHER INFORMATION: peptide)
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; Sequence 6, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
              Bard, Frederique
   APPLICANT:
  APPLICANT:
              Vasquez, Nicki
              Yednock, Ted
  APPLICANT:
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
   FILE REFERENCE: 15270J-004750UC
   CURRENT APPLICATION NUMBER: US/09/724,961
   CURRENT FILING DATE: 2000-11-28
   PRIOR APPLICATION NUMBER: US 09/580,015
   PRIOR FILING DATE: 2000-05-26
   PRIOR APPLICATION NUMBER: US 09/322,289
   PRIOR FILING DATE: 1999-05-28
   PRIOR APPLICATION NUMBER: US 09/201,430
   PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: WO PCT/US00/14810
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 PRIOR APPLICATION NUMBER: US 60/067,740
; PRIOR FILING DATE: 1997-12-02
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   OTHER INFORMATION: from AN1792 sequence (human Abeta42, beta-amyloid
   OTHER INFORMATION: peptide)
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US-09-724-961-7
; Sequence 7, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
 APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
  APPLICANT: Vasquez, Nicki
  APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
   FILE REFERENCE: 15270J-004750UC
  CURRENT APPLICATION NUMBER: US/09/724,961
   CURRENT FILING DATE: 2000-11-28
  PRIOR APPLICATION NUMBER: US 09/580,015
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   PRIOR FILING DATE: 1998-04-07
   PRIOR APPLICATION NUMBER: US 60/067,740
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    OTHER INFORMATION: peptide)
US-09-724-961-7
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US-09-724-961-8
; Sequence 8, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
; APPLICANT: Schenk, Dale B.
; APPLICANT: Bard, Frederique
; APPLICANT: Vasquez, Nicki
  APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
  FILE REFERENCE: 15270J-004750UC
 CURRENT APPLICATION NUMBER: US/09/724,961
; CURRENT FILING DATE: 2000-11-28
  PRIOR APPLICATION NUMBER: US 09/580,015
  PRIOR FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
  PRIOR APPLICATION NUMBER: US 09/201,430
   PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: WO PCT/US00/14810
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: US 60/080,970
  PRIOR FILING DATE: 1998-04-07
 PRIOR APPLICATION NUMBER: US 60/067,740
 PRIOR FILING DATE: 1997-12-02
  NUMBER OF SEQ ID NOS: 77
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    OTHER INFORMATION: from AN1792 sequence (human Abeta42, beta-amyloid
    OTHER INFORMATION: peptide)
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4 EFRH 7

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; Sequence 9, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
  APPLICANT: Vasquez, Nicki
  APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
  FILE REFERENCE: 15270J-004750UC
  CURRENT APPLICATION NUMBER: US/09/724,961
  CURRENT FILING DATE: 2000-11-28
  PRIOR APPLICATION NUMBER: US 09/580,015
  PRIOR FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
  PRIOR APPLICATION NUMBER: US 09/201,430
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: WO PCT/US00/14810
  PRIOR FILING DATE: 1998-11-30
; PRIOR APPLICATION NUMBER: US 60/080,970
; PRIOR FILING DATE: 1998-04-07
; PRIOR APPLICATION NUMBER: US 60/067,740
; PRIOR FILING DATE: 1997-12-02
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; Sequence 10, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
               Bard, Frederique
  APPLICANT:
               Vasquez, Nicki
; APPLICANT:
               Yednock, Ted
 APPLICANT:
; TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
; FILE REFERENCE: 15270J-004750UC
; CURRENT APPLICATION NUMBER: US/09/724,961
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CURRENT FILING DATE: 2000-11-28
  PRIOR APPLICATION NUMBER: US 09/580,015
  PRIOR FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
  PRIOR APPLICATION NUMBER: US 09/201,430
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: WO PCT/US00/14810
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: US 60/080,970
  PRIOR FILING DATE: 1998-04-07
  PRIOR APPLICATION NUMBER: US 60/067,740
  PRIOR FILING DATE: 1997-12-02
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   OTHER INFORMATION: peptide)
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RESULT 10
US-09-724-961-11
; Sequence 11, Application US/09724961
; Patent No. 6743427
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
  APPLICANT: Vasquez, Nicki
  APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
   FILE REFERENCE: 15270J-004750UC
  CURRENT APPLICATION NUMBER: US/09/724,961
  CURRENT FILING DATE: 2000-11-28
  PRIOR APPLICATION NUMBER: US 09/580,015
   PRIOR FILING DATE: 2000-05-26
   PRIOR APPLICATION NUMBER: US 09/322,289
   PRIOR FILING DATE: 1999-05-28
   PRIOR APPLICATION NUMBER: US 09/201,430
   PRIOR FILING DATE: 1998-11-30
   PRIOR APPLICATION NUMBER: WO PCT/US00/14810
  PRIOR FILING DATE: 1998-11-30
  PRIOR APPLICATION NUMBER: US 60/080,970
; PRIOR FILING DATE: 1998-04-07
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PRIOR APPLICATION NUMBER: US 60/067,740
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   OTHER INFORMATION: peptide)
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US-09-580-018-5
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; Patent No. 6761888
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
  APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
  FILE REFERENCE: 15270J-004760US
  CURRENT APPLICATION NUMBER: US/09/580,018
  CURRENT FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
  NUMBER OF SEQ ID NOS: 77
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    OTHER INFORMATION: from AN1792 sequence (human Abeta42, beta-amyloid
    OTHER INFORMATION: peptide)
US-09-580-018-5
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US-09-580-018-6
; Sequence 6, Application US/09580018
; Patent No. 6761888
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
  APPLICANT: Yednock, Ted
  TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
  FILE REFERENCE: 15270J-004760US
  CURRENT APPLICATION NUMBER: US/09/580,018
  CURRENT FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
  NUMBER OF SEQ ID NOS: 77
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    OTHER INFORMATION: peptide)
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Db
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; Sequence 7, Application US/09580018
; Patent No. 6761888
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
  APPLICANT: Yednock, Ted
   TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
   FILE REFERENCE: 15270J-004760US
   CURRENT APPLICATION NUMBER: US/09/580,018
  CURRENT FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
   PRIOR FILING DATE: 1999-05-28
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    TYPE: PRT
    ORGANISM: Artificial Sequence
    FEATURE:
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   OTHER INFORMATION: peptide)
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US-09-580-018-8
; Sequence 8, Application US/09580018
; Patent No. 6761888
; GENERAL INFORMATION:
  APPLICANT: Schenk, Dale B.
  APPLICANT: Bard, Frederique
              Yednock, Ted
  APPLICANT:
   TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
   FILE REFERENCE: 15270J-004760US
   CURRENT APPLICATION NUMBER: US/09/580,018
  CURRENT FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
  NUMBER OF SEQ ID NOS: 77
   SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 8
   LENGTH: 10
    TYPE: PRT
    ORGANISM: Artificial Sequence
    OTHER INFORMATION: Description of Artificial Sequence: 10-mer peptide
    OTHER INFORMATION: from AN1792 sequence (human Abeta42, beta-amyloid
    OTHER INFORMATION: peptide)
US-09-580-018-8
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US-09-580-018-9
; Sequence 9, Application US/09580018
; Patent No. 6761888
; GENERAL INFORMATION:
; APPLICANT: Schenk, Dale B.
; APPLICANT: Bard, Frederique
; APPLICANT: Yednock, Ted
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TITLE OF INVENTION: Prevention and Treatment of Amyloidogenic Disease
  FILE REFERENCE: 15270J-004760US
  CURRENT APPLICATION NUMBER: US/09/580,018
  CURRENT FILING DATE: 2000-05-26
  PRIOR APPLICATION NUMBER: US 09/322,289
  PRIOR FILING DATE: 1999-05-28
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; SEQ ID NO 9
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   ORGANISM: Artificial Sequence
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   OTHER INFORMATION: from AN1792 sequence (human Abeta42, beta-amyloid
   OTHER INFORMATION: peptide)
US-09-580-018-9
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Search completed: November 19, 2004, 17:00:23 Job time: 5.68085 secs

## GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: November 19, 2004, 16:35:52; Search time 4.08511 Seconds

(without alignments)

94.212 Million cell updates/sec

Title: US-09-830-954A-1

Perfect score: 24

Sequence: 1 EFRH 4

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: PIR\_79:\*

1: pir1:\*

2: pir2:\*

3: pir3:\*

4: pir4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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2	24	100.0	42	2	PN0512	beta-amyloid prote
3	24	100.0	52	2	C91112	hypothetical prote
4	24	100.0	57	2	A60045	Alzheimer's diseas
5	24	100.0	57	2	F60045	Alzheimer's diseas
6	24	100.0	57	2	D60045	Alzheimer's diseas
7	24	100.0	57	2	E60045	Alzheimer's diseas
8	24	100.0	57	2	G60045	Alzheimer's diseas
9	24	100.0	57	2	B60045	Alzheimer's diseas
10	24	100.0	57	2	B89981	truncated transpos
11	24	100.0	82	2	PQ0438	Alzheimer's diseas
12	24	100.0	84	2	G96025	hypothetical prote
13	24	100.0	89	2	C82331	hypothetical prote

14	24	100.0	91	2	T16095	hypothetical prote
15	24	100.0	94	2	B86195	hypothetical prote
16	24	100.0	97	1	RCBP22	abc2 protein - pha
17	24	100.0	97	2	н84901	hypothetical prote
18	24	100.0	106	2	G72059	conserved hypothet
19	24	100.0	106	2	D86563	CT466 hypothetical
20	24	100.0	116	2	B89964	truncated transpos
21	24	100.0	123	2	G95878	probable TRm2011-2
22	24	100.0	132	2	JQ0737	RnpA protein - Mic
23	24	100.0	133	2	AH2580	PTS system, IIA co
24	24	100.0	133	2	F97362	PTS enzyme IIAB, m
25	24	100.0	134	2	B86720	conserved hypothet
26	24	100.0	136	2	B56338	phospholipase A2 (
27	24	100.0	136	2	A87681	conserved hypothet
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31	24	100.0	141	2	D85605	unknown in ISEc8 [
32	24	100.0	141	2	A99803	hypothetical prote
33	24	100.0	141	2	E85611	unknown protein in
34	24	100.0	143	2	F75475	3-dehydroquinate d
35	24	100.0	145	2	AD2740	3-dehydroquinate d
36	24	100.0	148	2	B69960	3-dehydroquinate d
37	24	100.0	152	2	D75367	hypothetical prote
38	24	100.0	155	2	AC1187	B. subtilis YdcK p
39	2.4	100.0	155	2	AB1545	B. subtilis YdcK p
40	24	100.0	156	2	T02166	cysteine proteinas
41	24	100.0	162	2	B97521	3-dehydroquinate d
42	24	100.0	166	2	A28127	myosin light chain
43	24	100.0	167	2	Т34963	hypothetical prote
44	24	100.0	176	2	н72201	conserved hypothet
45	24	100.0	176	2	D95322	hypothetical prote

#### ALIGNMENTS

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RESULT 1
A48544
neuropeptide F - brown garden snail
C; Species: Helix aspersa (brown garden snail)
C; Date: 19-Nov-1993 #sequence_revision 18-Nov-1994 #text_change 09-Jul-2004
C; Accession: A48544
R; Leung, P.S.; Shaw, C.; Maule, A.G.; Thim, L.; Johnston, C.F.; Irvine, G.B.
Regul. Pept. 41, 71-81, 1992
A; Title: The primary structure of neuropeptide F (NPF) from the garden snail,
Helix aspersa.
A; Reference number: A48544; MUID:93087780; PMID:1472263
A; Accession: A48544
A; Status: preliminary
A; Molecule type: protein
A; Residues: 1-39 < LEU>
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A; Note: sequence extracted from NCBI backbone (NCBIP:120485)

A; Cross-references: UNIPROT: P41321

A; Experimental source: circumesophageal ganglia

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Best Local Similarity 100.0%; Pred. No. 22;
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Qу
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Db
RESULT 2
PN0512
beta-amyloid protein - guinea pig (fragment)
C; Species: Cavia porcellus (guinea pig)
C;Date: 31-Dec-1993 #sequence revision 31-Dec-1993 #text change 09-Jul-2004
C; Accession: PN0512
R; Shimohigashi, Y.; Matsumoto, H.; Takano, Y.; Saito, R.; Iwata, T.; Kamiya, H.;
Biochem. Biophys. Res. Commun. 193, 624-630, 1993
A; Title: Receptor-mediated specific biological activity of a beta-amyloid
protein fragment for NK-1 substance p receptors.
A; Reference number: PN0512; MUID: 93290653; PMID: 7685598
A; Accession: PN0512
A; Molecule type: protein
A; Residues: 1-42 <SHI>
A; Cross-references: UNIPROT: Q7M088
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
proteinase inhibitor homology
C; Keywords: alternative splicing; amyloid
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Qу
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C91112
hypothetical protein ECs3867 [imported] - Escherichia coli (strain 0157:H7,
substrain RIMD 0509952)
C; Species: Escherichia coli
C;Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text change 09-Jul-2004
C; Accession: C91112
R; Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.;
Han, C.G.; Ohtsubo, E.; Nakayama, K.; Murata, T.; Tanaka, M.; Tobe, T.; Iida,
T.; Takami, H.; Honda, T.; Sasakawa, C.; Ogasawara, N.; Yasunaga, T.; Kuhara,
S.; Shiba, T.; Hattori, M.; Shinagawa, H.
DNA Res. 8, 11-22, 2001
A; Title: Complete genome sequence of enterohemorrhagic Escherichia coli 0157:H7
and genomic comparison with a laboratory strain K-12.
A; Reference number: A99629; MUID:21156231; PMID:11258796
A; Accession: C91112
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-52 <HAY>
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A; Cross-references: UNIPROT: Q8X2N9; GB: BA000007; PIDN: BAB37290.1; PID: g13363339;
GSPDB:GN00154
A; Experimental source: strain O157:H7, substrain RIMD 0509952
C; Genetics:
A; Gene: ECs3867
  Query Match
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A60045
Alzheimer's disease amyloid beta/A4 protein precursor - dog (fragment)
C; Species: Canis lupus familiaris (dog)
C; Date: 01-Dec-1992 #sequence revision 01-Dec-1992 #text change 28-Jul-1995
C; Accession: A60045
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A; Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
A; Reference number: A60045; MUID: 92017079; PMID: 1656157
A:Accession: A60045
A; Molecule type: mRNA
A; Residues: 1-57 < JOH>
A; Cross-references: EMBL: X56125
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
proteinase inhibitor homology
C; Keywords: alternative splicing; Alzheimer's disease; amyloid; brain
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Qу
              8 EFRH 11
Db
RESULT 5
Alzheimer's disease amyloid beta/A4 protein precursor - pig (fragment)
C; Species: Sus scrofa domestica (domestic pig)
C; Date: 01-Dec-1992 #sequence revision 01-Dec-1992 #text change 13-Aug-1999
C: Accession: F60045
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A; Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
A; Reference number: A60045; MUID: 92017079; PMID: 1656157
A; Accession: F60045
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A; Residues: 1-57 < JOH>
A;Cross-references: EMBL:X56127; NID:g1895; PIDN:CAA39592.1; PID:g1896
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
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C; Keywords: alternative splicing; Alzheimer's disease; amyloid; brain
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  Matches
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Qу
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Db
RESULT 6
D60045
Alzheimer's disease amyloid beta/A4 protein precursor - bovine (fragment)
C; Species: Bos primigenius taurus (cattle)
C;Date: 01-Dec-1992 #sequence revision 01-Dec-1992 #text change 28-Jul-1995
C; Accession: D60045
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A; Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
A; Reference number: A60045; MUID: 92017079; PMID: 1656157
A; Accession: D60045
A; Molecule type: mRNA
A; Residues: 1-57 < JOH>
A; Cross-references: EMBL: X56124
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
proteinase inhibitor homology
C; Keywords: alternative splicing; Alzheimer's disease; amyloid; brain
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Qy
              8 EFRH 11
Db
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E60045
Alzheimer's disease amyloid beta/A4 protein precursor - sheep (fragment)
C; Species: Ovis sp. (sheep)
C;Date: 01-Dec-1992 #sequence revision 01-Dec-1992 #text change 28-Jul-1995
C; Accession: E60045
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A; Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
A; Reference number: A60045; MUID: 92017079; PMID: 1656157
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A; Accession: E60045
A; Molecule type: mRNA
A; Residues: 1-57 < JOH>
A; Cross-references: EMBL: X56130
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
proteinase inhibitor homology
C; Keywords: alternative splicing; Alzheimer's disease; amyloid; brain
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Qy
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G60045
Alzheimer's disease amyloid beta/A4 protein precursor - guinea pig (fragment)
C; Species: Cavia porcellus (quinea pig)
C;Date: 01-Dec-1992 #sequence revision 01-Dec-1992 #text change 28-Jul-1995
C; Accession: G60045
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A; Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
A; Reference number: A60045; MUID: 92017079; PMID: 1656157
A; Accession: G60045
A; Molecule type: mRNA
A; Residues: 1-57 < JOH>
A; Cross-references: EMBL: X56126
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
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C; Keywords: alternative splicing; Alzheimer's disease; amyloid; brain
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Qу
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            8 EFRH 11
Db
RESULT 9
B60045
Alzheimer's disease amyloid beta/A4 protein precursor - polar bear (fragment)
C; Species: Ursus maritimus (polar bear)
C;Date: 01-Dec-1992 #sequence revision 01-Dec-1992 #text change 09-Jul-2004
C; Accession: B60045
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A; Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
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A; Reference number: A60045; MUID: 92017079; PMID: 1656157
A; Accession: B60045
A; Molecule type: mRNA
A; Residues: 1-57 < JOH>
A; Cross-references: UNIPROT: Q29149; EMBL: X56128; NID: q2165; PIDN: CAA39593.1;
PID:q2166
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
proteinase inhibitor homology
C; Keywords: alternative splicing; Alzheimer's disease; amyloid; brain
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Qу
              1111
Db
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RESULT 10
B89981
truncated transposase [imported] - Staphylococcus aureus (strain N315)
C; Species: Staphylococcus aureus
C; Date: 10-May-2001 #sequence revision 10-May-2001 #text change 09-Jul-2004
C; Accession: B89981
R; Kuroda, M.; Ohta, T.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui,
L.; Oguchi, A.; Aoki, K.; Nagai, Y.; Lian, J.; Ito, T.; Kanamori, M.; Matsumaru,
H.; Maruyama, A.; Murakami, H.; Hosoyama, A.; Mizutani-Ui, Y.; Kobayashi, N.;
Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K.; Hirakawa, H.; Kuhara, S.; Goto,
S.; Yabuzaki, J.; Kanehisa, M.; Yamashita, A.; Oshima, K.; Furuya, K.; Yoshino,
C.; Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiramatsu, K.
Lancet 357, 1225-1240, 2001
A; Title: Whole genome sequencing of meticillin-resistant Stapylococcus aureus.
A; Reference number: A89758; MUID:21311952; PMID:11418146
A; Accession: B89981
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-57 < KUR>
A; Cross-references: UNIPROT: Q99SW5; GB: BA000018; PID: q13701716; PIDN: BAB43009.1;
GSPDB:GN00149
A; Experimental source: strain N315
C; Genetics:
A; Gene: truncated-tnp
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PO0438
Alzheimer's disease amyloid A4 protein precursor - rabbit (fragment)
C; Species: Oryctolagus cuniculus (domestic rabbit)
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C;Date: 30-Sep-1993 #sequence revision 19-Oct-1995 #text change 19-Oct-1995
C; Accession: PQ0438; C60045
R; Davidson, J.S.; West, R.L.; Kotikalapudi, P.; Maroun, L.E.
Biochem. Biophys. Res. Commun. 188, 905-911, 1992
A; Title: Sequence and methylation in the beta/A4 region of the rabbit amyloid
precursor protein gene.
A; Reference number: PQ0438; MUID: 93075180; PMID: 1445331
A; Accession: PQ0438
A; Molecule type: DNA
A; Residues: 1-82 < DAV>
A; Cross-references: GB:M83558; GB:M83657
R; Johnstone, E.M.; Chaney, M.O.; Norris, F.H.; Pascual, R.; Little, S.P.
Brain Res. Mol. Brain Res. 10, 299-305, 1991
A:Title: Conservation of the sequence of the Alzheimer's disease amyloid peptide
in dog, polar bear and five other mammals by cross-species polymerase chain
reaction analysis.
A; Reference number: A60045; MUID: 92017079; PMID: 1656157
A; Accession: C60045
A; Molecule type: mRNA
A; Residues: 12-68 < JOH>
A; Cross-references: EMBL:X56129
C; Superfamily: Alzheimer's disease amyloid beta protein; animal Kunitz-type
proteinase inhibitor homology
C; Keywords: alternative splicing; Alzheimer's disease; amyloid; Down's syndrome
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Qу
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Db
RESULT 12
G96025
hypothetical protein SMb20779 [imported] - Sinorhizobium meliloti (strain 1021)
magaplasmid pSymB
C; Species: Sinorhizobium meliloti
C;Date: 24-Aug-2001 #sequence revision 24-Aug-2001 #text change 09-Jul-2004
C; Accession: G96025
R; Finan, T.M.; Weidner, S.; Wong, K.; Buhrmester, J.; Chain, P.; Vorholter,
F.J.; Hernandez-Lucas, I.; Becker, A.; Cowie, A.; Gouzy, J.; Golding, B.;
Puhler, A.
Proc. Natl. Acad. Sci. U.S.A. 98, 9889-9894, 2001
A; Title: The complete sequence of the 1,683-kb pSymB megaplasmid from the N2-
fixing endosymbiont Sinorhizobium meliloti.
A; Reference number: A95842; MUID: 21396508; PMID: 11481431
A; Accession: G96025
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-84 < KUR>
A;Cross-references: UNIPROT:Q92TN7; GB:AL591985; PIDN:CAC49871.1; PID:g15141359;
GSPDB:GN00167
A; Experimental source: strain 1021, megaplasmid pSymB
R; Galibert, F.; Finan, T.M.; Long, S.R.; Puhler, A.; Abola, P.; Ampe, F.;
Barloy-Hubler, F.; Barnett, M.J.; Becker, A.; Boistard, P.; Bothe, G.; Boutry,
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M.; Bowser, L.; Buhrmester, J.; Cadieu, E.; Capela, D.; Chain, P.; Cowie, A.;
Davis, R.W.; Dreano, S.; Federspiel, N.A.; Fisher, R.F.; Gloux, S.; Godrie, T.;
Goffeau, A.; Golding, B.; Gouzy, J.; Gurjal, M.; Hernandez-Lucas, I.; Hong, A.;
Huizar, L.; Hyman, R.W.; Jones, T.
Science 293, 668-672, 2001
A; Authors: Kahn, D.; Kahn, M.L.; Kalman, S.; Keating, D.H.; Kiss, E.; Komp, C.;
Lelaure, V.; Masuy, D.; Palm, C.; Peck, M.C.; Pohl, T.M.; Portetelle, D.;
Purnelle, B.; Ramsperger, U.; Surzycki, R.; Thebault, P.; Vandenbol, M.;
Vorholter, F.J.; Weidner, S.; Wells, D.H.; Wong, K.; Yeh, K.C.; Batut, J.
A; Title: The composite genome of the legume symbiont Sinorhizobium meliloti.
A; Reference number: A96039; MUID:21368234; PMID:11474104
A; Contents: annotation
C; Genetics:
A; Gene: SMb20779
A; Genome: plasmid
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C;Date: 18-Aug-2000 #sequence revision 20-Aug-2000 #text change 09-Jul-2004
C; Accession: C82331
R; Heidelberg, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.;
Dodson, R.J.; Haft, D.H.; Hickey, E.K.; Peterson, J.D.; Umayam, L.A.; Gill,
S.R.; Nelson, K.E.; Read, T.D.; Tettelin, H.; Richardson, D.; Ermolaeva, M.D.;
Vamathevan, J.; Bass, S.; Qin, H.; Dragoi, I.; Sellers, P.; McDonald, L.;
Utterback, T.; Fleishmann, R.D.; Nierman, W.C.; White, O.; Salzberg, S.L.;
Smith, H.O.; Colwell, R.R.; Mekalanos, J.J.; Venter, J.C.; Fraser, C.M.
Nature 406, 477-483, 2000
A; Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio
cholerae.
A; Reference number: A82035; MUID: 20406833; PMID: 10952301
A; Accession: C82331
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-89 <HEI>
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C; Accession: T16095
R; Leimbach, D.
submitted to the EMBL Data Library, June 1995
A; Description: The sequence of C. elegans cosmid F18E9.
A; Reference number: Z18460
A; Accession: T16095
A; Status: preliminary; translated from GB/EMBL/DDBJ
A; Molecule type: DNA
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C; Accession: B86195
R; Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.;
Alonso, J.; Altaf, H.; Araujo, R.; Bowman, C.L.; Brooks, S.Y.; Buehler, E.;
Chan, A.; Chao, Q.; Chen, H.; Cheuk, R.F.; Chin, C.W.; Chung, M.K.; Conn, L.;
Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar, K.; Dunn, P.; Etgu, P.;
Feldblyum, T.V.; Feng, J.; Fong, B.; Fujii, C.Y.; Gill, J.E.; Goldsmith, A.D.;
Haas, B.; Hansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000
A; Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.;
Kim, C.J.; Koo, H.L.; Kremenetskaia, I.; Kurtz, D.B.; Kwan, A.; Lam, B.; Langin-
Hooper, S.; Lee, A.; Lee, J.M.; Lenz, C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu,
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S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Marziali, A.; Militscher, J.; Miranda, M.; Nguyen, M.; Nierman, W.C.; Osborne, B.I.; Pai, G.; Peterson, J.; Pham, P.K.;

A; Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon, L.J.; Tambunga, G.; Toriumi, M.J.; Town, C.D.; Utterback, T.; van Aken, S.; Vaysberg, M.; Vysotskaia, V.S.; Walker, M.; Wu, D.; Yu, G.; Fraser, C.M.;

Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.

Venter, J.C.; Davis, R.W.

A; Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis. A; Reference number: A86141; MUID:21016719; PMID:11130712 A; Accession: B86195 A;Status: preliminary A; Molecule type: DNA A; Residues: 1-94 <STO> A; Cross-references: UNIPROT: Q9LNE5; GB: AE005172; NID: g8810463; PIDN: AAF80124.1; GSPDB:GN00141 C; Genetics: A; Map position: 1 Query Match 100.0%; Score 24; DB 2; Length 94; Best Local Similarity 100.0%; Pred. No. 55; Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0; 1 EFRH 4 Qу

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Job time: 6.08511 secs

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## GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

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(without alignments)

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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2	24	100.0	4	9	US-09-975-932-8	Sequence 8, Appli
3	24	100.0	4	14	US-10-084-380A-8	Sequence 8, Appli
4	24	100.0	4	14	US-10-162-889-1	Sequence 1, Appli
. 5	24	100.0	4	15	US-10-384-788-1	Sequence 1, Appli
6	24	100.0	4	15	US-10-618-856-1	Sequence 1, Appli
7	24	100.0	6	9	US-09-808-037-7	Sequence 7, Appli
8	24	100.0	6	9	US-09-975-932-6	Sequence 6, Appli
9	24	100.0	6	14	US-10-084-380A-6	Sequence 6, Appli
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11	24	100.0	6	15	US-10-384-788-7	Sequence 7, Appli
12	24	100.0	6	15	US-10-618-856-7	Sequence 7, Appli
13	24	100.0	6	16	US-10-622-087-75	Sequence 75, Appl
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# ALIGNMENTS

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<sup>;</sup> Sequence 1, Application US/09808037 ; Patent No. US20020052311A1

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; GENERAL INFORMATION:
 APPLICANT: SOLOMON, Beka
  APPLICANT: HANAN, Eilat
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE TREATMENT AND/OR
DIAGNOSIS OF
  TITLE OF INVENTION: NEUROLOGICAL DISEASES AND DISORDERS
  FILE REFERENCE: SOLOMON=2D
  CURRENT APPLICATION NUMBER: US/09/808,037
  CURRENT FILING DATE: 2001-03-15
  PRIOR APPLICATION NUMBER: 09/629,971
  PRIOR FILING DATE: 2000-07-31
  PRIOR APPLICATION NUMBER: US 09/473,653
  PRIOR FILING DATE: 1999-12-29
  PRIOR APPLICATION NUMBER: US 60/152,417
  PRIOR FILING DATE: 1999-09-03
  NUMBER OF SEQ ID NOS: 33
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   TYPE: PRT
   ORGANISM: Artificial Sequence
   FEATURE:
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; Publication No. US20020086847A1
; GENERAL INFORMATION:
; APPLICANT: CHAIN, Daniel G.
  TITLE OF INVENTION: RECOMBINANT ANTIBODIES SPECIFIC FOR BETA-AMYLOID ENDS,
  TITLE OF INVENTION: DNA ENCODING AND METHODS OF USE THEREOF
  FILE REFERENCE: CHAIN1B
  CURRENT APPLICATION NUMBER: US/09/975,932
   CURRENT FILING DATE: 2001-10-15
  PRIOR APPLICATION NUMBER: 09/402,820
; PRIOR FILING DATE: 1999-10-12
  PRIOR APPLICATION NUMBER: PCT/US98/06900
  PRIOR FILING DATE: 1998-04-09
  PRIOR APPLICATION NUMBER: 60/041,850
  PRIOR FILING DATE: 1997-04-09
   NUMBER OF SEQ ID NOS: 8
   SOFTWARE: PatentIn Ver. 2.0
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    TYPE: PRT
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; Sequence 8, Application US/10084380A
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; GENERAL INFORMATION:
; APPLICANT: Mindset Biopharmaceutical Inc.
; APPLICANT: Chain, Daniel G.
; TITLE OF INVENTION: specific antibodies to amyloid beta peptide,
pharmaceutical compositions
; TITLE OF INVENTION: and methods of use thereof
   FILE REFERENCE: P-4815-US1
 CURRENT APPLICATION NUMBER: US/10/084,380A
   CURRENT FILING DATE: 2002-02-28
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   APPLICANT: HANAN, Eilat
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; TITLE OF INVENTION: USEFUL IN DIAGNOSING
; TITLE OF INVENTION: AND/OR TREATING OR PREVENTING PLAQUE FORMING DISEASES
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 APPLICANT: FRENKEL, Dan
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR TREATING A PLAQUE-FORMING
DISEASE
; FILE REFERENCE: SOLOMON=2D.2
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  CURRENT FILING DATE: 2003-03-11
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; PRIOR APPLICATION NUMBER: 60/152,417
; PRIOR FILING DATE: 1999-09-03
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; Publication No. US20040052766A1
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  APPLICANT: SOLOMON, Beka
  APPLICANT: FRENKEL, Dan
  TITLE OF INVENTION: IMMUNIZATION AGAINST AMYLOID PLAQUES USING DISPLAY
TECHNOLOGY
; FILE REFERENCE: SOLOMON=2A
  CURRENT APPLICATION NUMBER: US/10/618,856
  CURRENT FILING DATE: 2003-07-15
  PRIOR APPLICATION NUMBER: US/09/473,653A
  PRIOR FILING DATE: 1999-12-29
  PRIOR APPLICATION NUMBER: US 60/152,417
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; APPLICANT: SOLOMON, Beka
 APPLICANT: HANAN, Eilat
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DIAGNOSIS OF
; TITLE OF INVENTION: NEUROLOGICAL DISEASES AND DISORDERS
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; GENERAL INFORMATION:
; APPLICANT: CHAIN, Daniel G.
 TITLE OF INVENTION: RECOMBINANT ANTIBODIES SPECIFIC FOR BETA-AMYLOID ENDS,
; TITLE OF INVENTION: DNA ENCODING AND METHODS OF USE THEREOF
  FILE REFERENCE: CHAIN1B
  CURRENT APPLICATION NUMBER: US/09/975,932
; CURRENT FILING DATE: 2001-10-15
  PRIOR APPLICATION NUMBER: 09/402,820
; PRIOR FILING DATE: 1999-10-12
 PRIOR APPLICATION NUMBER: PCT/US98/06900
 PRIOR FILING DATE: 1998-04-09
; PRIOR APPLICATION NUMBER: 60/041,850
 PRIOR FILING DATE: 1997-04-09
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; Sequence 6, Application US/10084380A
; Publication No. US20030073655A1
; GENERAL INFORMATION:
; APPLICANT: Mindset Biopharmaceutical Inc.
; APPLICANT: Chain, Daniel G.
; TITLE OF INVENTION: specific antibodies to amyloid beta peptide,
pharmaceutical compositions
; TITLE OF INVENTION: and methods of use thereof
  FILE REFERENCE: P-4815-US1
  CURRENT APPLICATION NUMBER: US/10/084,380A
; CURRENT FILING DATE: 2002-02-28
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; PRIOR APPLICATION NUMBER: 09/402,820
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Qу
              \perp
            3 EFRH 6
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US-10-162-889-7
; Sequence 7, Application US/10162889
; Publication No. US20030077252A1
; GENERAL INFORMATION:
; APPLICANT: SOLOMON, Beka
; APPLICANT: HANAN, Eilat
  TITLE OF INVENTION: AGENTS AND COMPOSITIONS AND METHODS UTILIZING SAME
; TITLE OF INVENTION: USEFUL IN DIAGNOSING
; TITLE OF INVENTION: AND/OR TREATING OR PREVENTING PLAQUE FORMING DISEASES
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FILE REFERENCE: SOLOMON=2B
  CURRENT APPLICATION NUMBER: US/10/162,889
  CURRENT FILING DATE: 2002-06-06
  PRIOR APPLICATION NUMBER: US/09/629,971
  PRIOR FILING DATE: 2000-07-31
  PRIOR APPLICATION NUMBER: US 09/473,653
  PRIOR FILING DATE: 1999-12-29
  PRIOR APPLICATION NUMBER: US 60/152,417
  PRIOR FILING DATE: 1999-09-03
  NUMBER OF SEQ ID NOS: 29
   SOFTWARE: PatentIn version 3.0
; SEQ ID NO 7
   LENGTH: 6
    TYPE: PRT
   ORGANISM: Artificial Sequence
    FEATURE:
    OTHER INFORMATION: synthetic peptide
US-10-162-889-7
                          100.0%; Score 24; DB 14; Length 6;
  Query Match
  Best Local Similarity 100.0%; Pred. No. 1.4e+06;
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  Matches
           1 EFRH 4
Qу
              3 EFRH 6
Db
RESULT 11
US-10-384-788-7
; Sequence 7, Application US/10384788
; Publication No. US20040013647A1
; GENERAL INFORMATION:
; APPLICANT: SOLOMON, Beka
   APPLICANT: FRENKEL, Dan
   TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR TREATING A PLAQUE-FORMING
DISEASE
   FILE REFERENCE: SOLOMON=2D.2
   CURRENT APPLICATION NUMBER: US/10/384,788
   CURRENT FILING DATE: 2003-03-11
   PRIOR APPLICATION NUMBER: 60/371,735
   PRIOR FILING DATE: 2002-04-12
   PRIOR APPLICATION NUMBER: 09/808,037
  PRIOR FILING DATE: 2001-03-15
  PRIOR APPLICATION NUMBER: 09/830,954
   PRIOR FILING DATE: 2001-06-22
   PRIOR APPLICATION NUMBER: 10/162,889
   PRIOR FILING DATE: 2002-06-06
  PRIOR APPLICATION NUMBER: 09/473,653
   PRIOR FILING DATE: 1999-12-29
   PRIOR APPLICATION NUMBER: 09/629,971
   PRIOR FILING DATE: 2000-07-31
  PRIOR APPLICATION NUMBER: 60/152,417
   PRIOR FILING DATE: 1999-09-03
   PRIOR APPLICATION NUMBER: PCT/IL00/00518
   PRIOR FILING DATE: 2000-08-31
   NUMBER OF SEQ ID NOS: 33
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SOFTWARE: PatentIn version 3.2
; SEQ ID NO 7
   LENGTH: 6
   TYPE: PRT
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: synthetic peptide
US-10-384-788-7
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Qу
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US-10-618-856-7
; Sequence 7, Application US/10618856
; Publication No. US20040052766A1
; GENERAL INFORMATION:
; APPLICANT: SOLOMON, Beka
; APPLICANT: FRENKEL, Dan
  TITLE OF INVENTION: IMMUNIZATION AGAINST AMYLOID PLAQUES USING DISPLAY
TECHNOLOGY
; FILE REFERENCE: SOLOMON=2A
  CURRENT APPLICATION NUMBER: US/10/618,856
  CURRENT FILING DATE: 2003-07-15
; PRIOR APPLICATION NUMBER: US/09/473,653A
  PRIOR FILING DATE: 1999-12-29
  PRIOR APPLICATION NUMBER: US 60/152,417
  PRIOR FILING DATE: 1999-09-03
  NUMBER OF SEQ ID NOS: 26
  SOFTWARE: PatentIn version 3.0
; SEQ ID NO 7
   LENGTH: 6
    TYPE: PRT
   ORGANISM: Artificial Sequence
    FEATURE:
    OTHER INFORMATION: synthetic peptide
US-10-618-856-7
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           4; Conservative 0; Mismatches 0;
                                                     Indels
                                                                0; Gaps
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Qy.
              \Box\Box\Box
            3 EFRH 6
RESULT 13
US-10-622-087-75
; Sequence 75, Application US/10622087
; Publication No. US20040141984A1
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; GENERAL INFORMATION:
  APPLICANT: Bachmann, Martin F
  APPLICANT: Tissot, Alain
  APPLICANT: Ortmann, Rainer
  APPLICANT: Luond, Rainer
  APPLICANT: Staufenbiel, Matthias
  APPLICANT: Frey, Peter
  TITLE OF INVENTION: Amyloid Beta 1-6 Antigen Arrays
  FILE REFERENCE: 1700.0350002
  CURRENT APPLICATION NUMBER: US/10/622,087
  CURRENT FILING DATE: 2003-07-18
  PRIOR APPLICATION NUMBER: US 60/396,639
  PRIOR FILING DATE: 2002-07-19
  PRIOR APPLICATION NUMBER: US 60/470,432
 PRIOR FILING DATE: 2003-05-15
 NUMBER OF SEQ ID NOS: 93
  SOFTWARE: PatentIn version 3.2
; SEQ ID NO 75
   LENGTH: 6
   TYPE: PRT
   ORGANISM: Homo sapiens
US-10-622-087-75
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Qy
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Db
           3 EFRH 6
RESULT 14
US-10-622-087-84
; Sequence 84, Application US/10622087
; Publication No. US20040141984A1
; GENERAL INFORMATION:
; APPLICANT: Bachmann, Martin F
  APPLICANT: Tissot, Alain
  APPLICANT: Ortmann, Rainer
  APPLICANT: Luond, Rainer
  APPLICANT: Staufenbiel, Matthias
  APPLICANT: Frey, Peter
; TITLE OF INVENTION: Amyloid Beta 1-6 Antigen Arrays
  FILE REFERENCE: 1700.0350002
  CURRENT APPLICATION NUMBER: US/10/622,087
  CURRENT FILING DATE: 2003-07-18
  PRIOR APPLICATION NUMBER: US 60/396,639
  PRIOR FILING DATE: 2002-07-19
  PRIOR APPLICATION NUMBER: US 60/470,432
  PRIOR FILING DATE: 2003-05-15
 NUMBER OF SEQ ID NOS: 93
  SOFTWARE: PatentIn version 3.2
; SEQ ID NO 84
   LENGTH: 6
   TYPE: PRT
   ORGANISM: homo sapiens
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Query Match
                        100.0%; Score 24; DB 16; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
          4; Conservative 0; Mismatches
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 Matches
                                              0; Indels
           1 EFRH 4
Qу
             -1111
           3 EFRH 6
Db
RESULT 15
US-10-622-087-85
; Sequence 85, Application US/10622087
; Publication No. US20040141984A1
; GENERAL INFORMATION:
; APPLICANT: Bachmann, Martin F
; APPLICANT: Tissot, Alain
; APPLICANT: Ortmann, Rainer
 APPLICANT: Luond, Rainer
  APPLICANT: Staufenbiel, Matthias
  APPLICANT: Frey, Peter
  TITLE OF INVENTION: Amyloid Beta 1-6 Antigen Arrays
 FILE REFERENCE: 1700.0350002
; CURRENT APPLICATION NUMBER: US/10/622,087
; CURRENT FILING DATE: 2003-07-18
; PRIOR APPLICATION NUMBER: US 60/396,639
; PRIOR FILING DATE: 2002-07-19
; PRIOR APPLICATION NUMBER: US 60/470,432
  PRIOR FILING DATE: 2003-05-15
; NUMBER OF SEQ ID NOS: 93
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 85
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   TYPE: PRT
   ORGANISM: Oryctolagus cuniculus
US-10-622-087-85
  Query Match
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  Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 4; Conservative 0; Mismatches
                                              0; Indels
                                                            0; Gaps
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Qу
           1 EFRH 4
             -1111
Db
           3 EFRH 6
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Search completed: November 19, 2004, 17:15:18

Job time : 16.6809 secs

### GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: November 19, 2004, 16:35:27; Search time 21.4468 Seconds

(without alignments)

107.312 Million cell updates/sec

Title: US-09-830-954A-1

Perfect score: 24

Sequence: 1 EFRH 4

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: UniProt 02:\*

1: uniprot\_sprot:\*
2: uniprot\_trembl:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

			¥					
Re	esult		Query					
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	1	24	100.0	25	2	Q7R843		plasmodium
	2	24	100.0	33	2	Q9UC33	Q9uc33	homo sapien
	3	24	100.0	35	2	Q8WZ99	Q8wz99	homo sapien
	4	24	100.0	38	2	Q8CM52	Q8cm52	staphylococ
	5	24	1,00.0	38	2	Q8CN66	Q8cn66	staphylococ
	6	24	100.0	38	2	Q8CNR0	Q8cnr0	staphylococ
	7	24	100.0	38	2	Q8CNT8	Q8cnt8	staphylococ
	8	24	100.0	38	2	Q8CP00	Q8cp00	staphylococ
	9	24	100.0	38	2	Q8CPT7	Q8cpt7	staphylococ
	10	24	100.0	38	2	Q8CPX2	Q8cpx2	staphylococ
	11	24	100.0	39	1	NPF HELAS	P41321	helix asper
	12	24	100.0	42	2	Q7M088	Q7m088	cavia porce
	13	24	100.0	43	2	Q7UT20	Q7ut20	rhodopirell
	14	24	100.0	45	2	Q6V7T6	Q6v7t6	burkholderi
	15	24	100.0	45	2	Q8CQ63	Q8cq63	staphylococ

16	24	100.0	45	2	AAQ54942	Aaq54942 burkholde
17	24	100.0	48	2	Q8CN63	Q8cn63 staphylococ
18	24	100.0	52	2	Q8X2N9	Q8x2n9 escherichia
19	24	100.0	55	2	Q7UGM8	Q7ugm8 rhodopirell
20	24	100.0	57	1	A4 URSMA	Q29149 ursus marit
21	24	100.0	57	2	Q8DGN4	Q8dgn4 synechococc
22	24	100.0	57	2	Q99SW5	Q99sw5 staphylococ
23	24	100.0	58	1	A4 CANFA	Q28280 canis famil
24	24	100.0	58	1	A4 RABIT	Q28748 oryctolagus
25	24	100.0	58	1	A4 SHEEP	Q28757 ovis aries
26	24	100.0	59	1	A4 BOVIN	Q28053 bos taurus
27	24	100.0	62	2	Q65802	Q65802 bovine vira
28	24	100.0	62	2	Q65804	Q65804 bovine vira
29	24	100.0	62	2	Q65805	Q65805 bovine vira
30	24	100.0	65	2	Q6IIT6	Q6iit6 drosophila
31	24	100.0	66	2	Q84Z36	Q84z36 oryza sativ
32	24	100.0	67	2	Q98LA2	Q981a2 rhizobium 1
33	24	100.0	68	2	014885	O14885 homo sapien
34	24	100.0	69	2	Q98MZ4	Q98mz4 rhizobium l
35	24	100.0	73	2	Q8GX82	Q8gx82 arabidopsis
36	24	100.0	76	2	Q87L89	Q87189 vibrio para
37	24	100.0	76	2	Q8DCK3	Q8dck3 vibrio vuln
38	24	100.0	77	2	Q6LM57	Q61m57 photobacter
39	24	100.0	77	2	CAG21620	Cag21620 photobact
40	24	100.0	80	2	Q8GR41	Q8gr41 enterococcu
41	24	100.0	80	2	Q7TCW4	Q7tcw4 untyped hum
42	24	100.0	84	2	Q92TN7	Q92tn7 rhizobium m
43	24	100.0	88	2	Q7MHA4	Q7mha4 vibrio vuln
44	24	100.0	89	2	Q9KUX5	Q9kux5 vibrio chol
45	24	100.0	90	2	Q8VZU1	Q8vzul arabidopsis

### ALIGNMENTS

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RESULT 1
Q7R843
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                                      PRT;
                                              25 AA.
ID
     Q7R843
AC
     Q7R843;
     01-MAR-2004 (TrEMBLrel. 26, Created)
01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DT
DT
     01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DT
     Hypothetical protein (Fragment).
DE
GN
     Name=PY07380;
     Plasmodium yoelii yoelii.
OS
     Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
OC
OX
     NCBI TaxID=73239;
RN
     [1]
RP
     SEQUENCE FROM N.A.
RC
     STRAIN=17XNL;
RX
     PubMed=12368865;
     Carlton J.M., Angiuoli S.V., Suh B.B., Kooij T.W., Pertea M.,
RA
     Silva J.C., Ermolaeva M.D., Allen J.E., Selengut J.D., Koo H.L.,
RA
     Peterson J.D., Pop M., Kosack D.S., Shumway M.F., Bidwell S.L.,
RA
     Shallom S.J., van Aken S.E., Riedmuller S.B., Feldblyum T.V.,
RA
     Cho J.K., Quackenbush J., Sedegah M., Shoaibi A., Cummings L.M.,
RA
     Florens L., Yates F.R. III, Raine J.D., Sinden R.E., Harris M.A.,
RA
```

```
Cunningham D.A., Preiser P.R., Bergman L.W., Vaidya A.B.,
RA
     van Lin L.H., Janse C.J., Waters A.P., Smith H.O., White O.R.,
RA
     Salzberg S.L., Venter J.C., Fraser C.M., Hoffman S.L., Gardner M.J.,
RA
     Carucci D.J.;
RA
     "Genome sequence and comparative analysis of the model rodent malaria
RT
     parasite Plasmodium yoelii yoelii.";
RT
     Nature 419:512-519(2002).
RL
     -!- CAUTION: The sequence shown here is derived from an
CC
         EMBL/GenBank/DDBJ whole genome shotgun (WGS) entry which is
CC
         preliminary data.
CC
     EMBL; AABL01002689; EAA19793.1; -.
DR
KW
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     NON TER
FT
                   1
                25 AA; 3303 MW; 1A5AB86BD78F4422 CRC64;
     SEOUENCE
SQ
                          100.0%; Score 24; DB 2; Length 25;
  Query Match
  Best Local Similarity
                          100.0%; Pred. No. 1.1e+02;
                                                                              0;
                                 0; Mismatches
                                                    0; Indels
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             4; Conservative
            1 EFRH 4
Qу
              1111
           10 EFRH 13
Db
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Q9UC33
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                 PRELIMINARY;
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                                            33 AA.
TD
AC
     Q9UC33;
     01-MAY-2000 (TrEMBLrel. 13, Created)
ידים
     01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DΤ
     01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DΤ
     Beta-amyloid peptide (Fragment).
DE
OS
     Homo sapiens (Human).
     Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC
     Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OC
OX
     NCBI TaxID=9606;
RN
     [1]
RP
     SEQUENCE.
     MEDLINE=93024877; PubMed=1406936;
RX
     Seubert P., Vigo-Pelfrey C., Esch F., Lee M., Dovey H., Davis D.,
RA
     Sinha S., Schlossmacher M., Whaley J., Swindlehurst C.;
RA
     "Isolation and quantification of soluble Alzheimer's beta-peptide from
RT
     biological fluids.";
RT
     Nature 359:325-327(1992).
RL
     GO; GO:0016021; C:integral to membrane; IEA.
DR
DR
     GO; GO:0005488; F:binding; IEA.
     InterPro; IPR001255; Beta-APP.
DR
     Pfam; PF03494; Beta-APP; 1.
DR
     PRINTS; PR00204; BETAAMYLOID.
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                33 AA; 3674 MW; B1DEFE2F4167ABD0 CRC64;
SO
                           100.0%; Score 24; DB 2; Length 33;
  Query Match
                           100.0%; Pred. No. 1.5e+02;
  Best Local Similarity
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                                                                               0;
             4; Conservative
                                 0; Mismatches
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            1 EFRH 4
Qy
              1111
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Q8WZ99
                  PRELIMINARY;
                                    PRT;
                                             35 AA.
ID
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AC
     01-MAR-2002 (TrEMBLrel. 20, Created)
DT
     01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT
     01-MAR-2002 (TrEMBLrel. 20, Last annotation update)
DT
     Amyloid protein (Fragment).
DE
     Name=APP;
GN
     Homo sapiens (Human).
OS
     Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC
     Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OC
     NCBI TaxID=9606;
OX
RN
     [1]
     SEQUENCE FROM N.A.
RP
     PubMed=15201367;
RX
     Wakutani Y., Watanabe K., Adachi Y., Wada-Isoe K., Urakami K.,
RA
     Ninomiya H., Saido TC., Hashimoto T., Iwatsubo T., Nakashima K.;
RA
     "Novel amyloid precursor protein gene missense mutation (D678N) in
RT
RT
     probable familial Alzheimer's disease.";
     J. Neurol. Neurosurg. Psychiatr. 75:1039-1042(2004).
RL
     EMBL; AB066441; BAB71958.2; -.
DR
     NON TER
                           1
FT
                   1
FT
     NON TER
                   35
                          35
SQ
     SEQUENCE
                 35 AA;
                         4084 MW;
                                   49D7D17289743B71 CRC64;
                           100.0%; Score 24; DB 2; Length 35;
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                                                                                 0;
              4; Conservative
                                  0; Mismatches 0;
                                                         Indels
                                                                    0;
                                                                        Gaps
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Qу
               \parallel \parallel \parallel \parallel
           19 EFRH 22
RESULT 4
Q8CM52
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ID
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                                             38 AA.
AC
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     01-MAR-2003 (TrEMBLrel. 23, Created)
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
     01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT
DE
     Truncated transposase.
     OrderedLocusNames=SE0257;
GN
     Staphylococcus epidermidis.
OS
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
     NCBI TaxID=1282;
OX
RN
RP
     SEQUENCE FROM N.A.
RC
     STRAIN=ATCC 12228;
RX
     PubMed=12950922;
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
```

```
"Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
    Mol. Microbiol. 49:1577-1593(2003).
RL
    EMBL; AE016744; AA003854.1; -.
DR
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KW
               38 AA; 4448 MW; A40B39C53421AD0E CRC64;
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                          100.0%; Pred. No. 1.7e+02;
  Best Local Similarity
                                                                             0;
            4; Conservative
                              0; Mismatches 0; Indels
                                                                 0; Gaps
 Matches
           1 EFRH 4
Qу
              30 EFRH 33
Db
RESULT 5
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                                   PRT:
                                           38 AA.
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                 PRELIMINARY;
ID
AC
     Q8CN66;
     01-MAR-2003 (TrEMBLrel. 23, Created)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DТ
     01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT
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DΕ
     OrderedLocusNames=SE1982;
GN
     Staphylococcus epidermidis.
os
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
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OX
RN
     [1]
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RP
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RC
     PubMed=12950922;
RX
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
     Mol. Microbiol. 49:1577-1593(2003).
RL
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KW
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     SEQUENCE 38 AA; 4458 MW; 8BDB217272BB9946 CRC64;
SQ
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  Query Match
                          100.0%; Pred. No. 1.7e+02;
  Best Local Similarity
                                0; Mismatches
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                                                  0; Indels
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  Matches
            4; Conservative
            1 EFRH 4
Qу
              1111
           30 EFRH 33
Db
RESULT 6
Q8CNR0
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                                   PRT;
                                            38 AA.
ID
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AC
     Q8CNR0;
     01-MAR-2003 (TrEMBLrel. 23, Created)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT
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01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT
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DE
     OrderedLocusNames=SE1539;
GN
     Staphylococcus epidermidis.
OS
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
    NCBI TaxID=1282;
OX
RN
     [1]
     SEQUENCE FROM N.A.
RP
     STRAIN=ATCC 12228;
RC
     PubMed=12950922;
RX
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RТ
     Mol. Microbiol. 49:1577-1593(2003).
DR
     EMBL; AE016749; AA005138.1; -.
ΚW
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                38 AA; 4434 MW; A415C75C6E85FC4B CRC64;
     SEQUENCE
SQ
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  Query Match
  Best Local Similarity 100.0%; Pred. No. 1.7e+02;
                                0; Mismatches
                                                                  0; Gaps
                                                                               0;
             4; Conservative
                                                  0; Indels
  Matches
            1 EFRH 4
Qу
              +1111
           30 EFRH 33
Db
RESULT 7
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                                    PRT:
                                            38 AA.
     Q8CNT8
ID
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AC
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DT
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DТ
     01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT
     Truncated transposase.
DΕ
     OrderedLocusNames=SE1454;
GN
     Staphylococcus epidermidis.
OS
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
     NCBI TaxID=1282;
OX
RN
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RP
     STRAIN=ATCC 12228;
RC
RX
     PubMed=12950922;
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
     Mol. Microbiol. 49:1577-1593(2003).
RL
     EMBL; AE016748; AA005053.1; -.
DR
KW
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                38 AA; 4404 MW; FE0B21726435EC91 CRC64;
SO
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                           100.0%;
  Query Match
                           100.0%; Pred. No. 1.7e+02;
  Best Local Similarity
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0; Indels
                                                                  0; Gaps
                                                                              0;
             4; Conservative
                                  0; Mismatches
  Matches
            1 EFRH 4
Qy
              1111
           30 EFRH 33
Db
RESULT 8
08CP00
                                    PRT;
                                            38 AA.
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AC
     Q8CP00;
     01-MAR-2003 (TrEMBLrel. 23, Created)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT
DE
     Truncated transposase.
GN
     OrderedLocusNames=SE1319;
     Staphylococcus epidermidis.
OS
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
OX
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RN
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RC
     PubMed=12950922;
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     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
     Mol. Microbiol. 49:1577-1593(2003).
RL
     EMBL; AE016748; AA004918.1; -.
DR
      Complete proteome.
KW
                 38 AA; 4495 MW; B1D109D5DC4F3B0E CRC64;
      SEQUENCE
SO
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   Query Match
  Best Local Similarity 100.0%; Pred. No. 1.7e+02;
                                                                               0;
                                                                       Gaps
                                 0; Mismatches
                                                   0; Indels
                                                                   0;
              4; Conservative
  Matches
             1 EFRH 4
 Qy
               30 EFRH 33
 Db
 RESULT 9
 Q8CPT7
                                    PRT:
                                             38 AA.
                  PRELIMINARY;
 ΙD
      Q8CPT7
      Q8CPT7;
 АC
      01-MAR-2003 (TrEMBLrel. 23, Created)
 DT
      01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT
      01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DT
      Truncated transposase.
 DE
      OrderedLocusNames=SE0668;
 GN
      Staphylococcus epidermidis.
 OS
      Bacteria; Firmicutes; Bacillales; Staphylococcus.
 OC
      NCBI TaxID=1282;
 OX
 RN
 ŔР
      SEOUENCE FROM N.A.
      STRAIN=ATCC 12228;
 RC
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RX
     PubMed=12950922;
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
    Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
    Mol. Microbiol. 49:1577-1593(2003).
RL
     EMBL; AE016746; AA004265.1; -.
DR
KW
     Complete proteome.
                38 AA; 4534 MW; A40B3F052E90E80E CRC64;
     SEQUENCE
SQ
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  Query Match
                          100.0%; Pred. No. 1.7e+02;
  Best Local Similarity
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                                0; Mismatches
                                                      Indels
                                                                  0; Gaps
                                                   0;
  Matches
             4; Conservative
         ~ 1 EFRH 4
Qу
              1111
           30 EFRH 33
Db
RESULT 10
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                                   PRT;
                                           38 AA.
                 PRELIMINARY;
ID
     Q8CPX2
     Q8CPX2;
AC
     01-MAR-2003 (TrEMBLrel. 23, Created)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DΤ
     01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DT
     Truncated transposase.
DE
     OrderedLocusNames=SE0590;
GN
     Staphylococcus epidermidis.
os
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
     NCBI TaxID=1282;
OX
RN
     [1]
     SEQUENCE FROM N.A.
RP
     STRAIN=ATCC 12228;
RC
     PubMed=12950922;
RX
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
     Mol. Microbiol. 49:1577-1593(2003).
RL
     EMBL; AE016745; AA004187.1; -.
DR
KW
     Complete proteome.
                        4395 MW; A40B39DF8421AD0E CRC64;
     SEQUENCE
               38 AA;
SO
                           100.0%; Score 24; DB 2; Length 38;
  Query Match
                          100.0%;
                                    Pred. No. 1.7e+02;
  Best Local Similarity
                                                                  0; Gaps
                                0; Mismatches
                                                    0;
                                                       Indels
             4; Conservative
            1 EFRH 4
Qу
               1111
           30 EFRH 33
Db
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RESULT 11 NPF HELAS

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PRT;
                                             39 AA.
     NPF HELAS
                    STANDARD;
ID
     P41321;
AC
     01-FEB-1995 (Rel. 31, Created)
DT
     01-FEB-1995 (Rel. 31, Last sequence update)
DT
     05-JUL-2004 (Rel. 44, Last annotation update)
DT
     Neuropeptide F (NPF).
DE
     Helix aspersa (Brown garden snail).
OS
     Eukaryota; Metazoa; Mollusca; Gastropoda; Pulmonata; Stylommatophora;
OC
     Sigmurethra; Helicoidea; Helicidae; Helix.
OC
     NCBI TaxID=6535;
OX
     [1]
RN
     SEQUENCE.
RP
     TISSUE=Circumoesophageal ganglion;
RC
     MEDLINE=93087780; PubMed=1472263;
RX
     Leung P.S., Shaw C., Maule A.G., Thim L., Johnston C.F., Irvine G.B.;
RA
     "The primary structure of neuropeptide F (NPF) from the garden snail,
RT
RT
     Helix aspersa.";
     Regul. Pept. 41:71-81(1992).
RL
     -!- FUNCTION: May have an important physiological role in
CC
         neuroregulation.
CC
     -!- SUBCELLULAR LOCATION: Secreted.
CC
     -!- TISSUE SPECIFICITY: Neuronal somata and fibers.
CC
     -!- SIMILARITY: Belongs to the NPY family.
CC
     PIR; A48544; A48544.
DR
     HSSP; P41967; 1K8V.
DR
     InterPro; IPR001955; Pancreatic hormn.
DR
     Pfam; PF00159; Hormone 3; 1.
DR
     PROSITE; PS00265; PANCREATIC_HORMONE_1; 1.
DR
     PROSITE; PS50276; PANCREATIC HORMONE 2; 1.
DR
     Amidation; Direct protein sequencing; Neuropeptide.
KW
                   39
                          39
                                    Phenylalanine amide.
FT
     MOD RES
     SEQUENCE
                 39 AA;
                         4855 MW;
                                   4B54AA7414CAAE33 CRC64;
SQ
                           100.0%; Score 24; DB 1; Length 39;
  Query Match
  Best Local Similarity
                           100.0%; Pred. No. 1.8e+02;
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                                                                    0;
                                                                        Gaps
  Matches
              4; Conservative
                                  0; Mismatches
                                                   .0; Indels
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Qy
               13 EFRH 16
Db
RESULT 12
O7M088
                                             42 AA.
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                                     PRT;
ID
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AC
      Q7M088;
      01-MAR-2004 (TrEMBLrel. 26, Created)
01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DT
DT
      01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DT
      Beta-amyloid protein (Fragment).
OS.
      Cavia porcellus (Guinea pig).
      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC
      Mammalia; Eutheria; Rodentia; Hystricognathi; Caviidae; Cavia.
OC
OX
      NCBI TaxID=10141;
RN
      [1]
      SEQUENCE.
 RP
      MEDLINE=93290653; PubMed=7685598;
 RX
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RA
     Shimohigashi Y., Matsumoto H., Takano Y., Saito R., Iwata T.,
RA
     Kamiya H., Ohno M.;
RT
     "Receptor-mediated specific biological activity of a beta-amyloid
RT
     protein fragment for NK-1 substance p receptors.";
RL
     Biochem. Biophys. Res. Commun. 193:624-630(1993).
DR
     PIR; PN0512; PN0512.
DR
     GO; GO:0016021; C:integral to membrane; IEA.
DR
     GO; GO:0005488; F:binding; IEA.
DR
     InterPro; IPR001255; Beta-APP.
     Pfam; PF03494; Beta-APP; 1.
DR
DR
     PRINTS; PR00204; BETAAMYLOID.
FT
     NON_TER
                   1
                          1
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FT
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                         42
SQ
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                42 AA; 4514 MW;
                                  3AC85563D7858C37 CRC64;
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                                                     Length 42;
 Best Local Similarity
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 Matches
             4; Conservative
                                 0; Mismatches
                                                                              0;
                                                    0; Indels
                                                                  0; Gaps
            1 EFRH 4
Qу
              1111
Db
            3 EFRH 6
RESULT 13
Q7UT20
ID
     Q7UT20
                 PRELIMINARY;
                                   PRT:
                                            43 AA.
AC
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DT
     01-OCT-2003 (TrEMBLrel. 25, Created)
     01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT
DT
     01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE
     Hypothetical protein.
GN
     OrderedLocusNames=RB4166;
OS
     Rhodopirellula baltica.
OC
     Bacteria; Planctomycetes; Planctomycetacia; Planctomycetales;
OC
     Planctomycetaceae; Pirellula.
OX
     NCBI TaxID=117;
RN
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     SEQUENCE FROM N.A.
RP
RC
     STRAIN=1;
RX
     MEDLINE=22735913; PubMed=12835416;
RA
     Gloeckner F.O., Kube M., Bauer M., Teeling H., Lombardot T.,
RA
     Ludwig W., Gade D., Beck A., Borzym K., Heitmann K., Rabus R.,
RA
     Schlesner H., Amann R., Reinhardt R.;
RT
     "Complete genome sequence of the marine planctomycete Pirellula sp.
RT
     strain 1.";
     Proc. Natl. Acad. Sci. U.S.A. 100:8298-8303(2003).
RL
DR
     EMBL; BX294140; CAD73621.1; -.
KW
     Complete proteome; Hypothetical protein.
SQ
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               43 AA; 5428 MW; 3106E49E67D19882 CRC64;
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                                  Score 24; DB 2; Length 43;
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                          100.0%;
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                                 0; Mismatches
                                                    0; Indels
            1 EFRH 4
Qу.
              1111
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RESULT 14
Q6V7T6
                                           45 AA.
                 PRELIMINARY;
                                   PRT;
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ID
    Q6V7T6;
AC
     05-JUL-2004 (TrEMBLrel. 27, Created)
DT
     05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
     05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
DT
     Gp6.
DE
     Burkholderia cepacia phage Bcep22.
OS
     Viruses; dsDNA viruses, no RNA stage; Caudovirales; Podoviridae.
OC
    NCBI TaxID=242527;
OX
RN
     [1]
     SEQUENCE FROM N.A.
RP
     Summer E.J., Cordova M., Parkinson B.C., Fuller A.C., Kitsopoulos K.,
RA
     Parks B., Rambo L., Rothwell S., Mebane L.M., Carlile T.M., No E.G.,
RA
     Gonzalez C.M., Young R.F.;
RA
     Submitted (JUL-2003) to the EMBL/GenBank/DDBJ databases.
RL
DR
     EMBL; AY349011; AAQ54942.1; -.
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SQ
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                                                  0; Indels
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            4; Conservative
            1 EFRH 4
Qу
              1111
            7 EFRH 10
Db
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                                    PRT;
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ID
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AC
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     01-MAR-2003 (TrEMBLrel. 23, Created)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT
     01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT
     Truncated transposase.
DE
     OrderedLocusNames=SE0355;
GN
     Staphylococcus epidermidis.
OS
     Bacteria; Firmicutes; Bacillales; Staphylococcus.
OC
     NCBI TaxID=1282;
OX
RN
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RP
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     PubMed=12950922;
     Zhang Y.-Q., Ren S.-X., Li H.-L., Wang Y.-X., Fu G., Yang J.,
RA
     Qin Z.-Q., Miao Y.-G., Wang W.-Y., Chen R.-S., Shen Y., Chen Z.,
RA
     Yuan Z.-H., Zhao G.-P., Qu D., Danchin A., Wen Y.-M.;
RA
     "Genome-based analysis of virulence genes in a non-biofilm-forming
RT
     Staphylococcus epidermidis strain (ATCC 12228).";
RT
     Mol. Microbiol. 49:1577-1593(2003).
RL
     EMBL; AE016745; AA003952.1; -.
DR
KW
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      SEQUENCE
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Query Match 100.0%; Score 24; DB 2; Length 45; Best Local Similarity 100.0%; Pred. No. 2e+02; Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Search completed: November 19, 2004, 16:58:25 Job time: 24.4468 secs